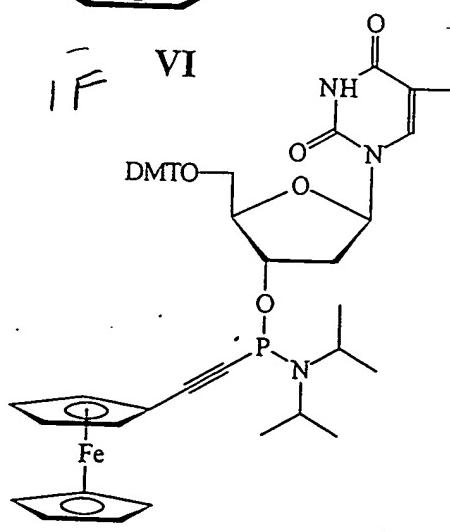
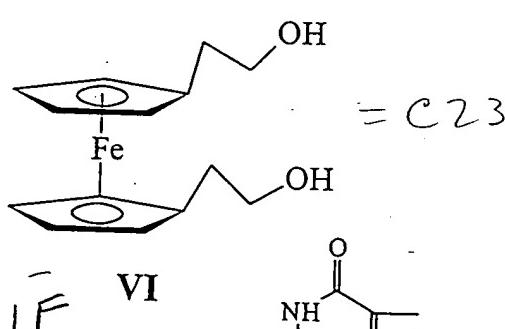
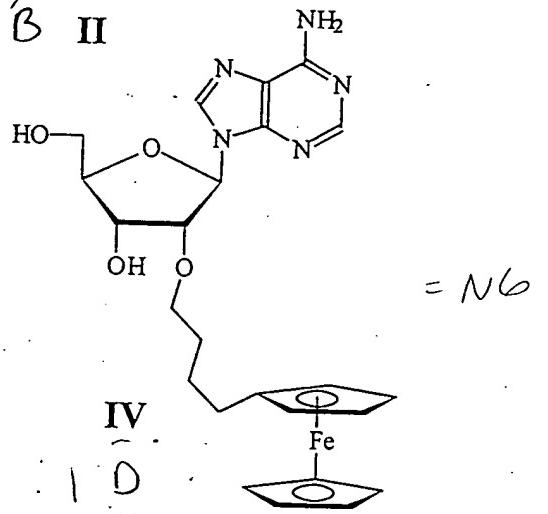
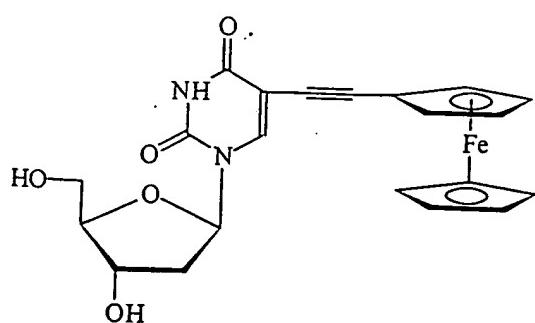
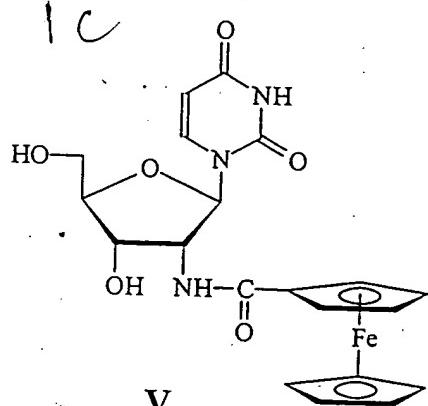
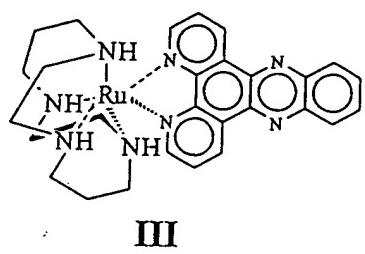
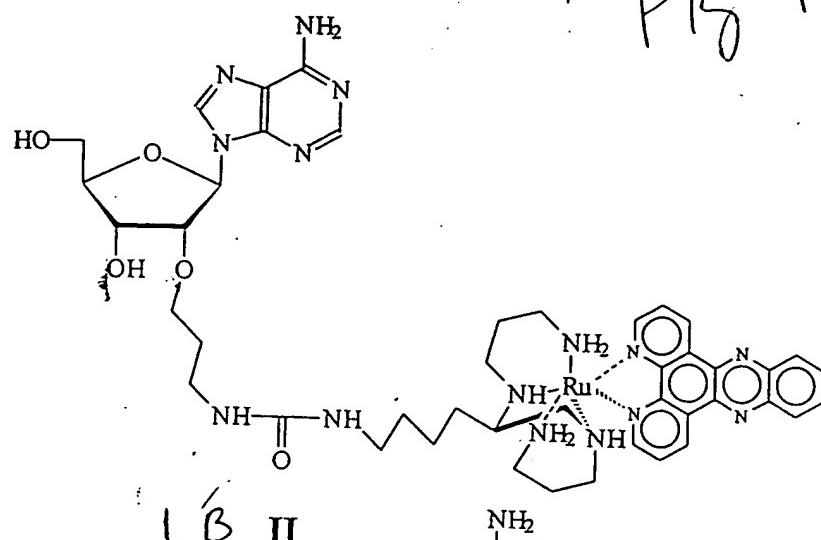
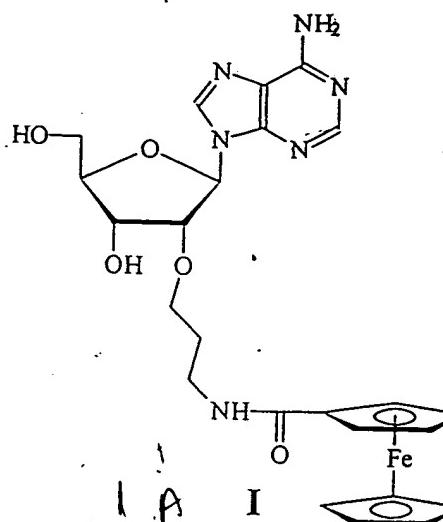


# List of Metal Complexes for Comparison

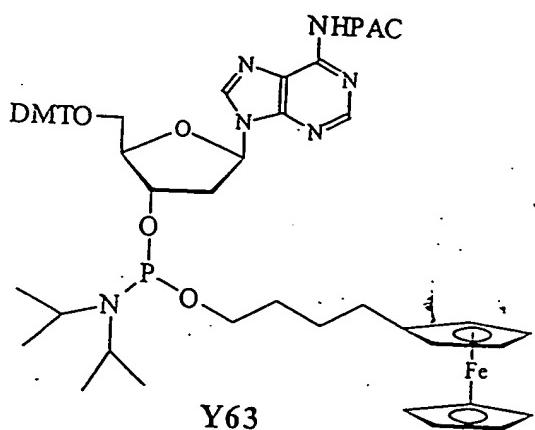
Fig 1



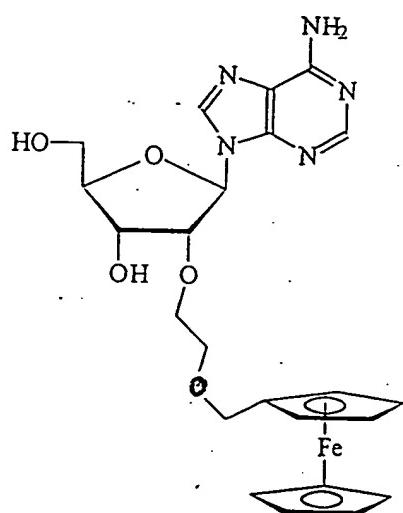
13H

118/98

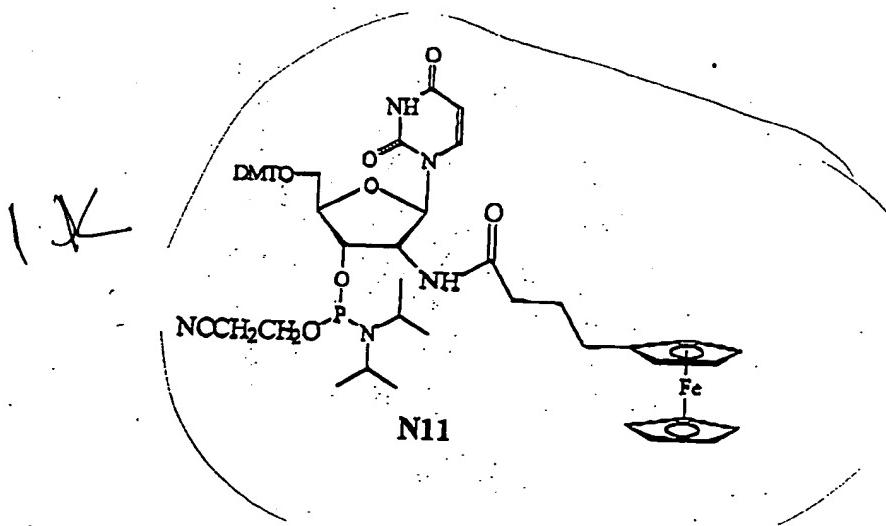
Fig 1  
(cont.)



11



1. J



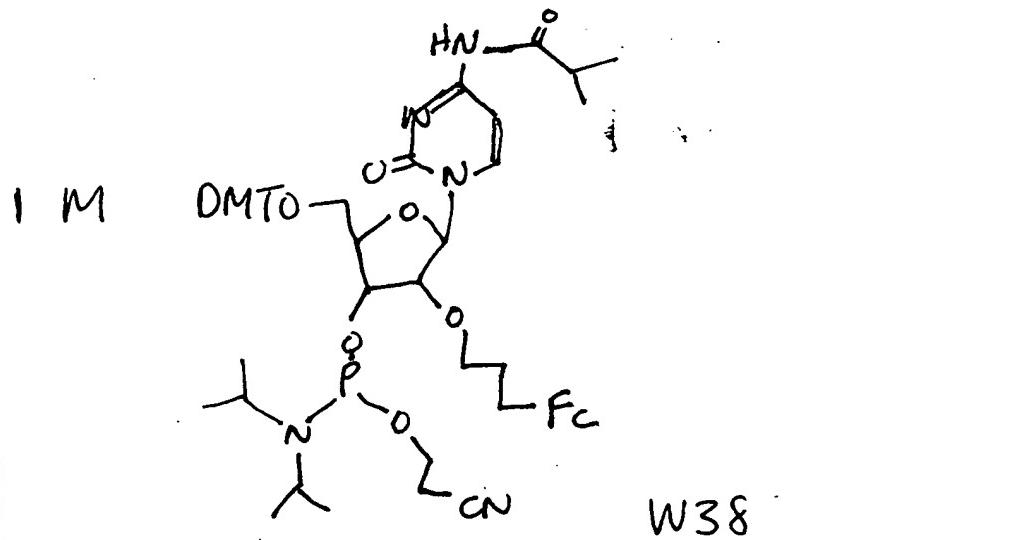
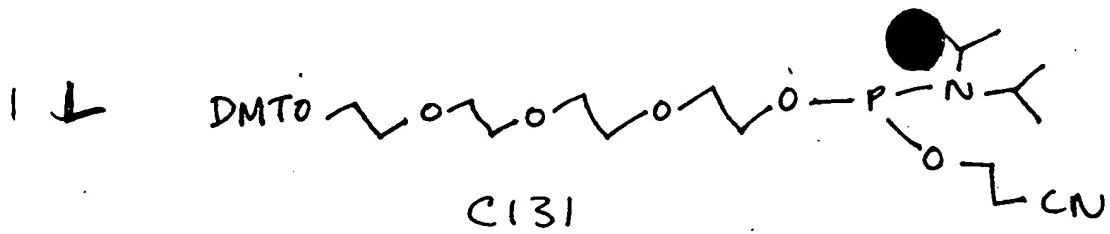
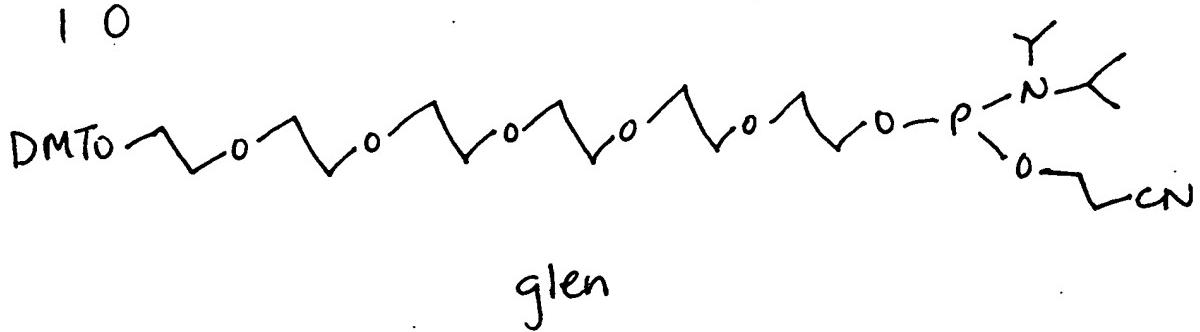
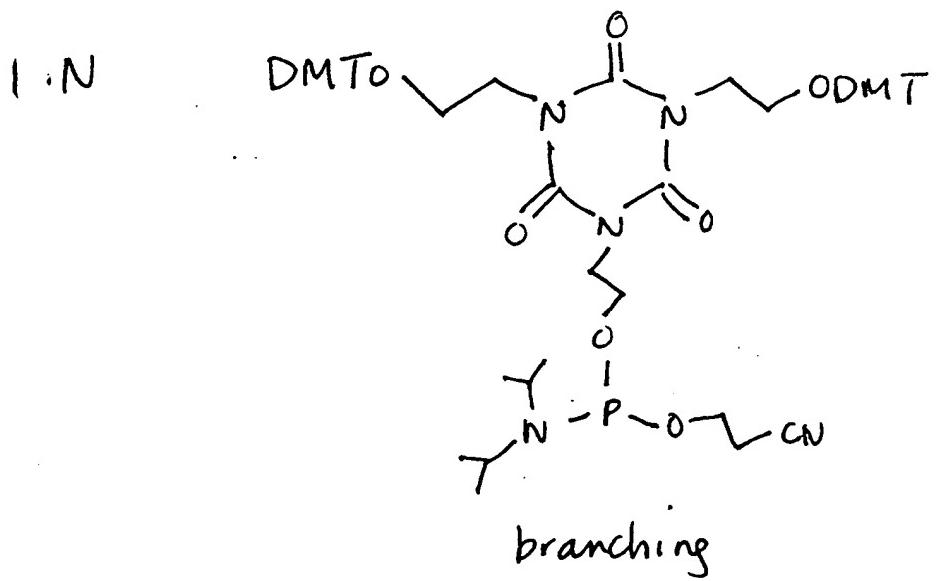


Fig 1  
(Cont.)



# Synthesis Scheme of Adenosine Ferrocene

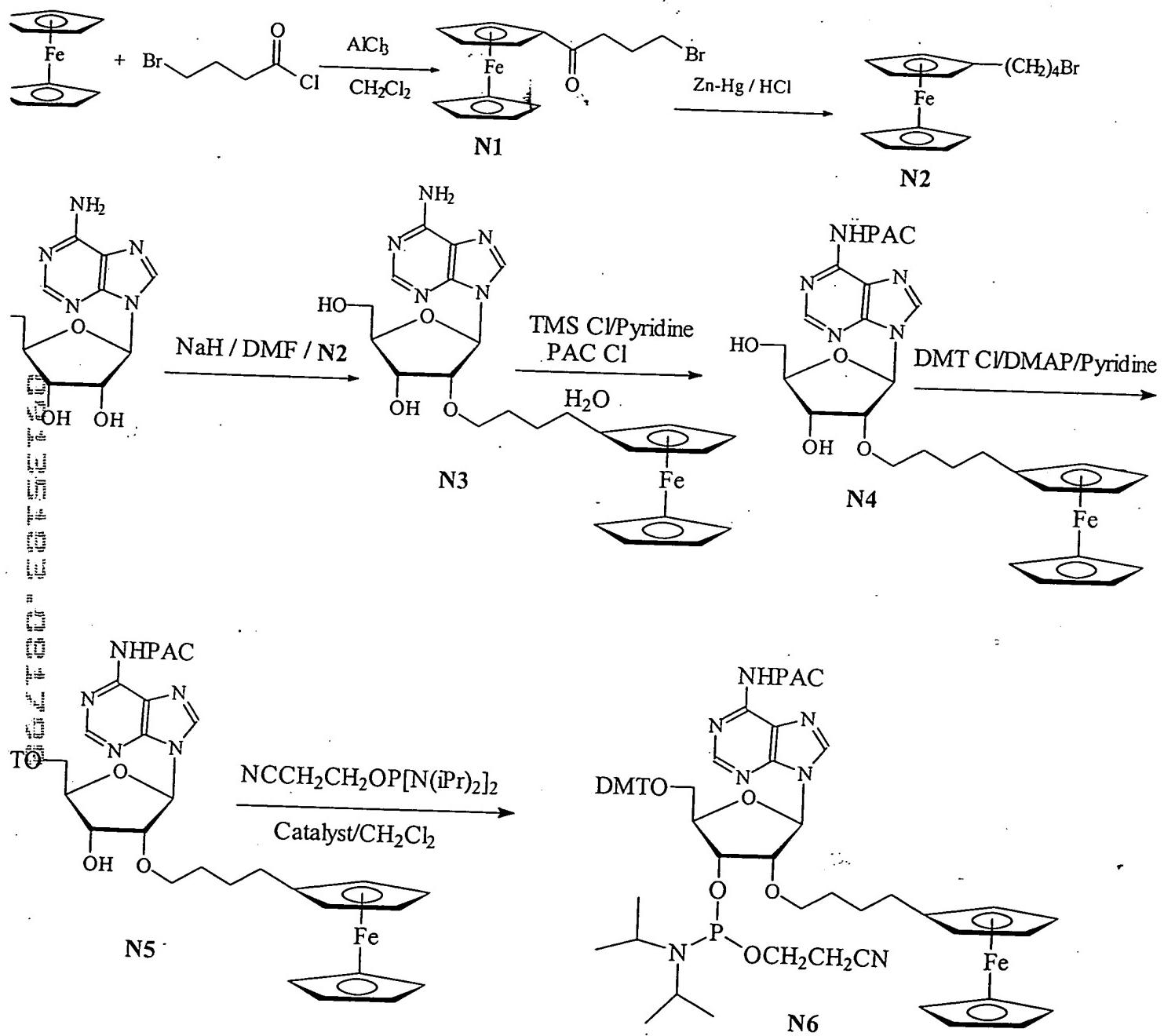


FIG.  
2

## Synthesis Scheme of Cytidine Ferrocene

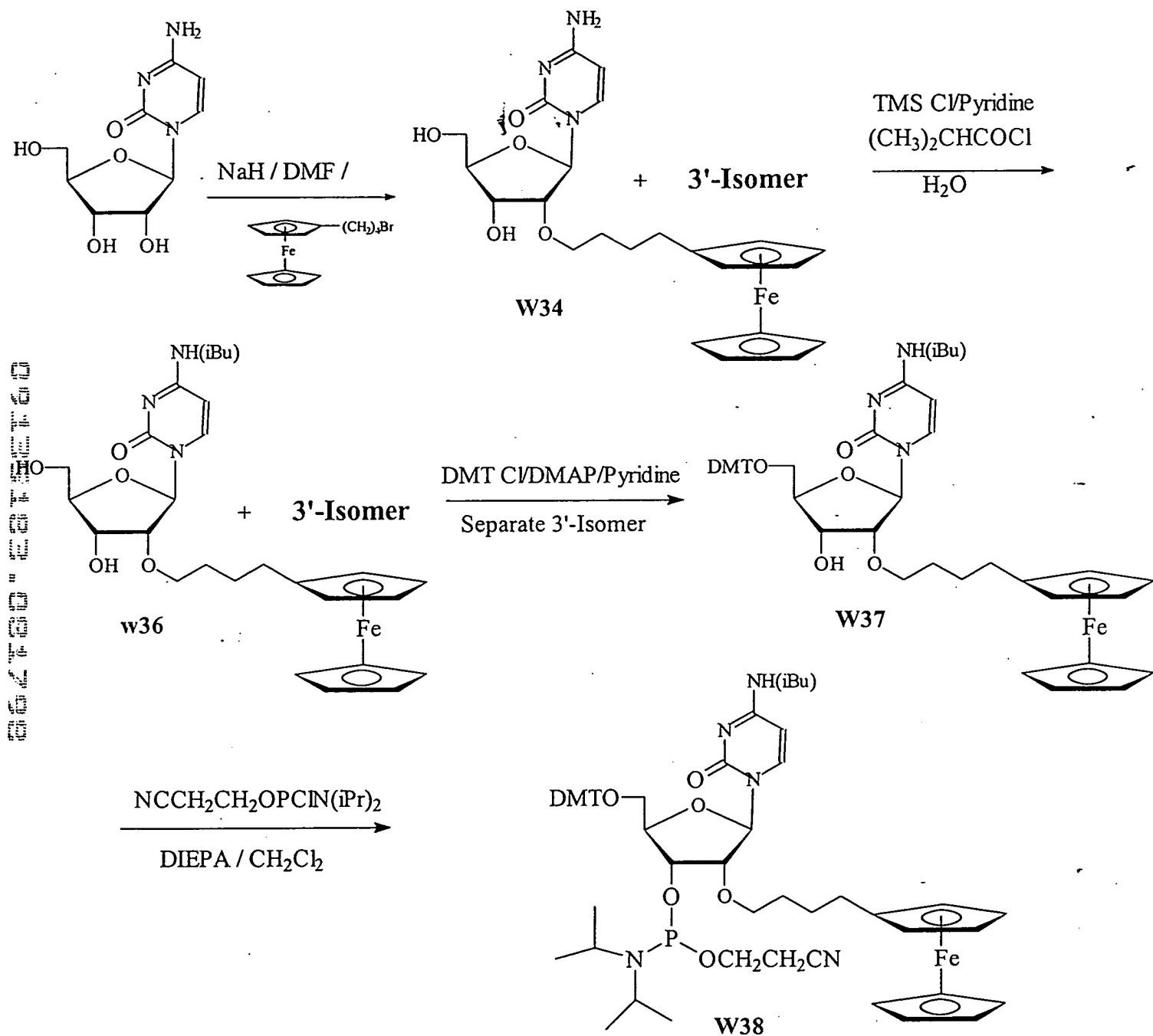


FIG.  
3

Synthesis Scheme of Y63

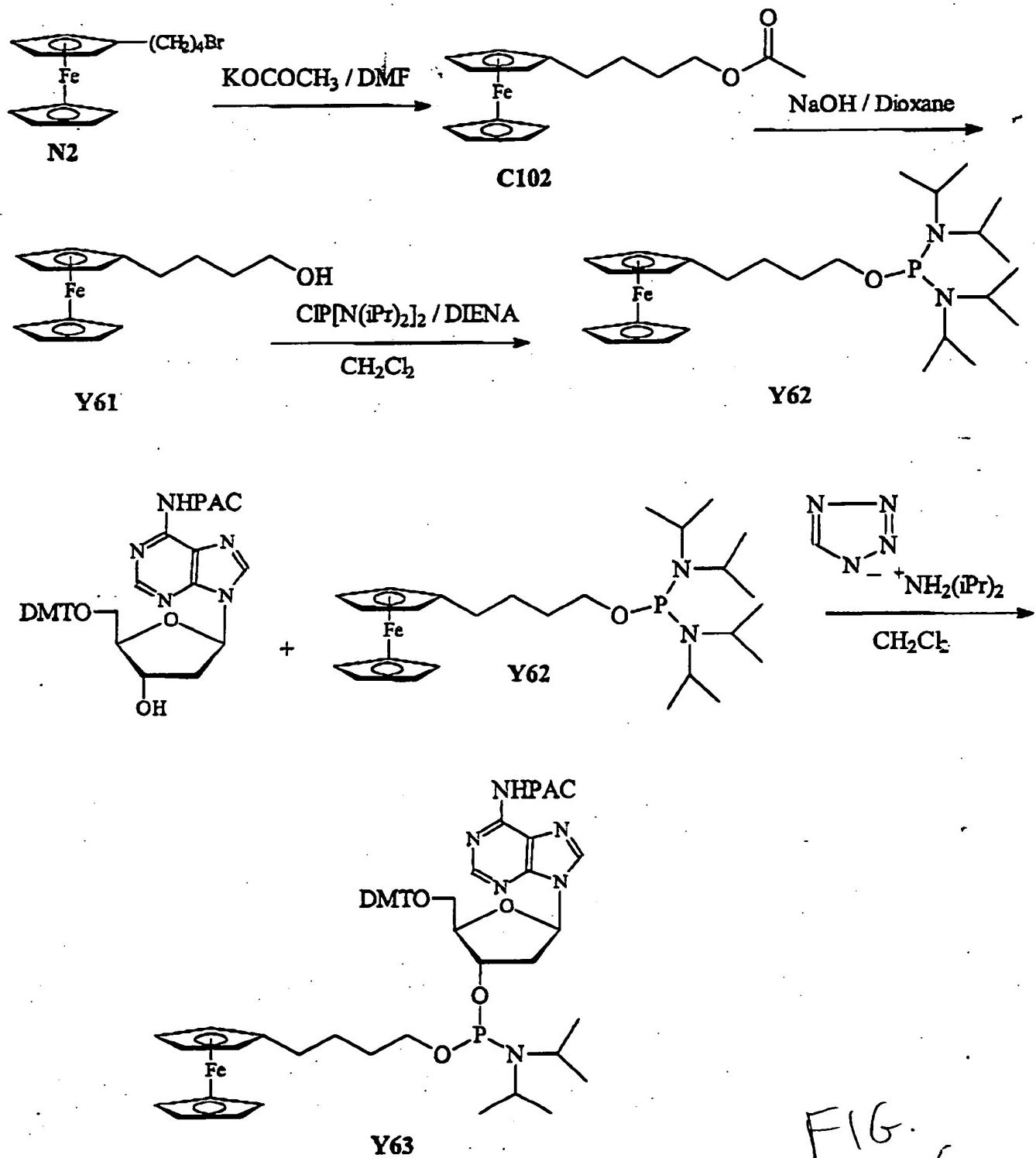


FIG.  
4

## Synthesis of Adenosine Ferrocene Triphosphate

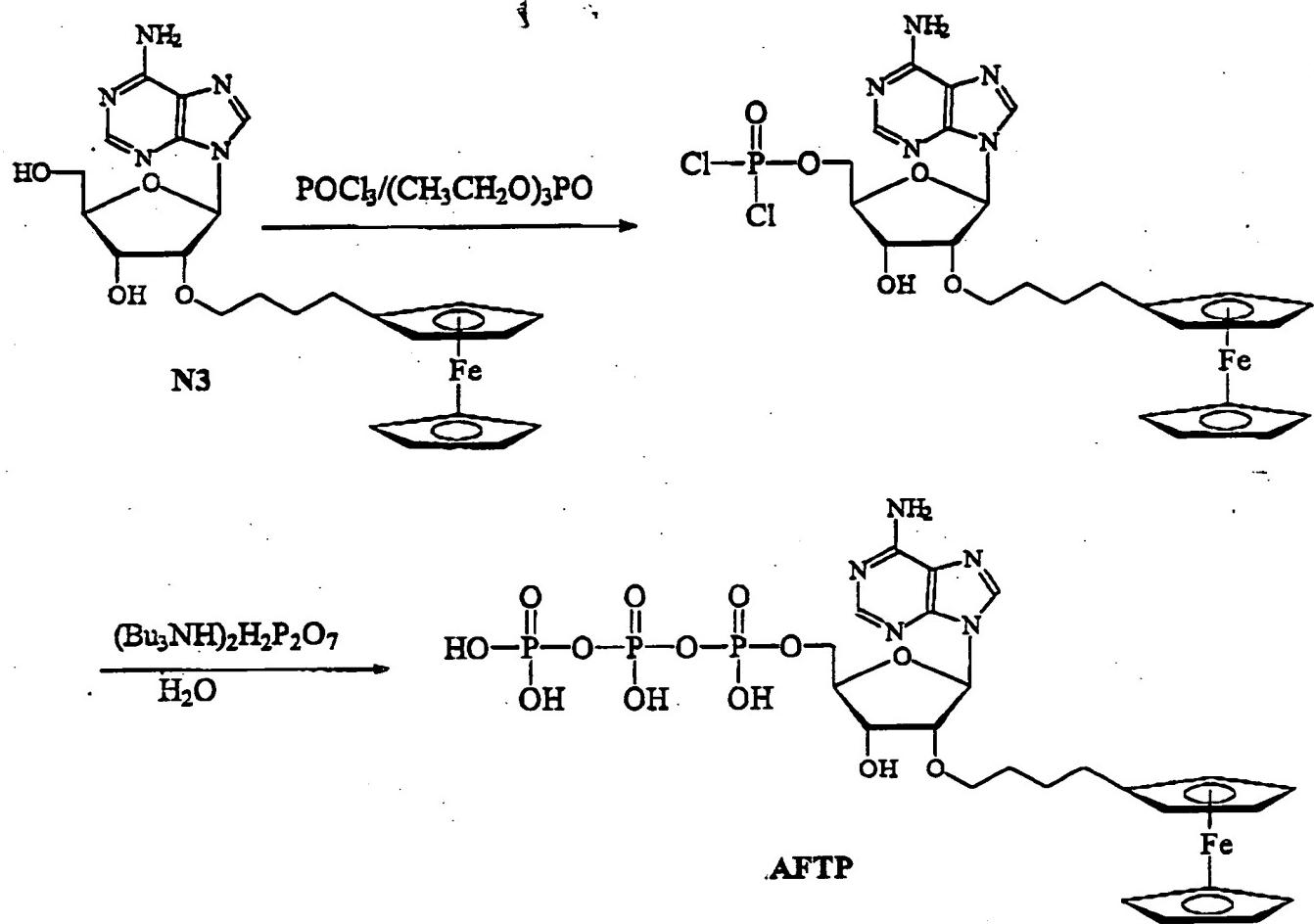


FIG. 5

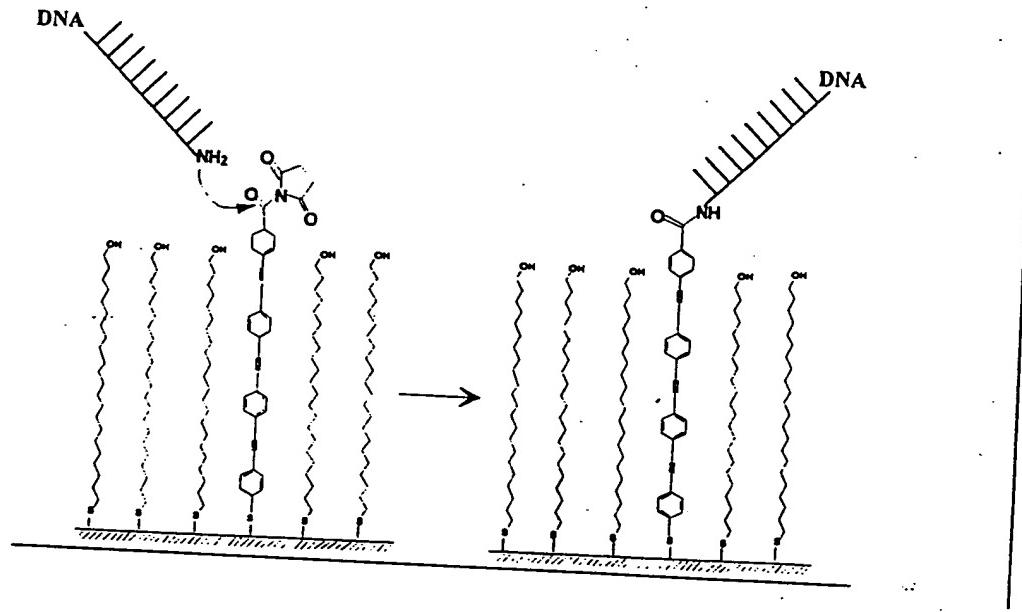


FIG.

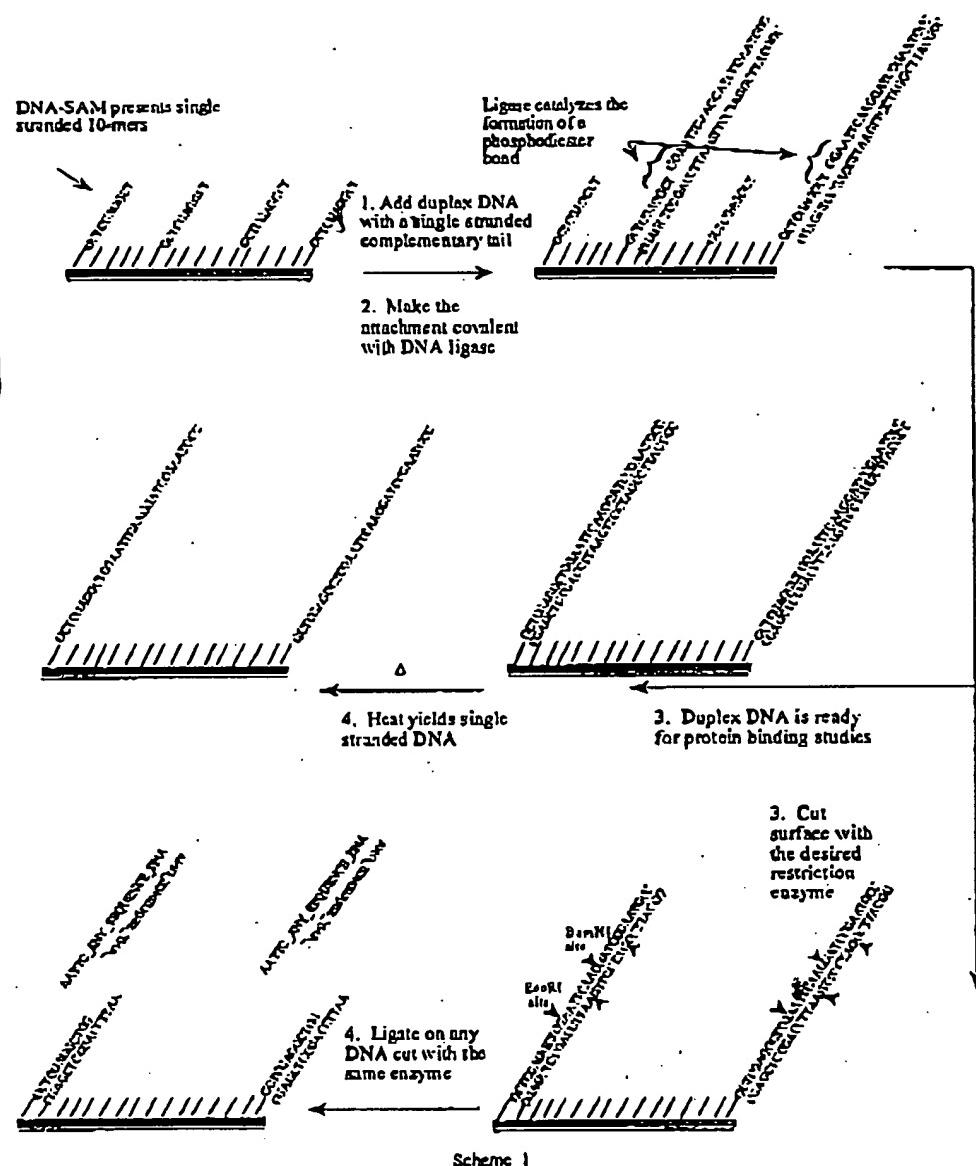
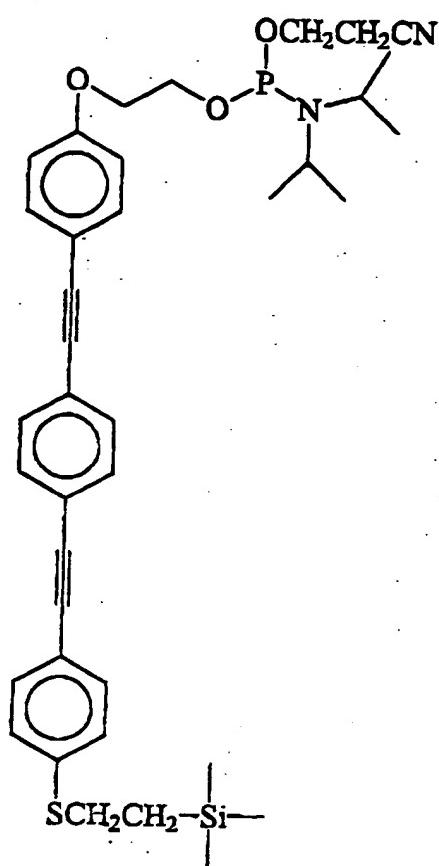
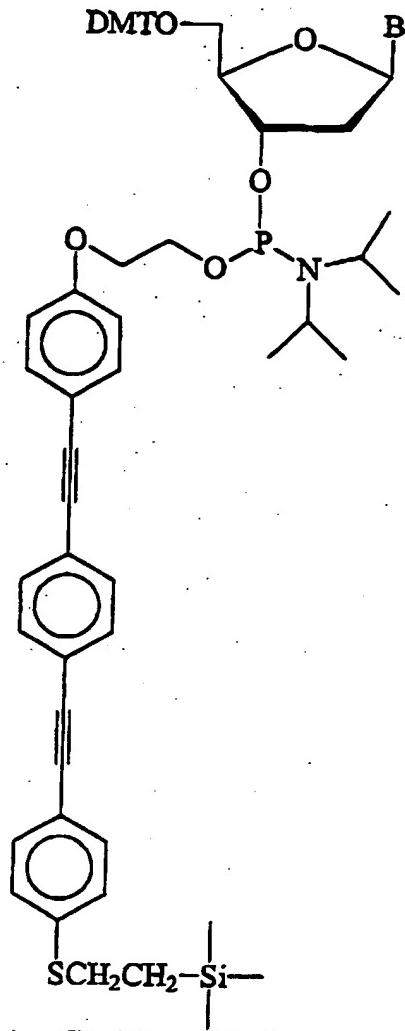


FIG. 7



5'-Attachment



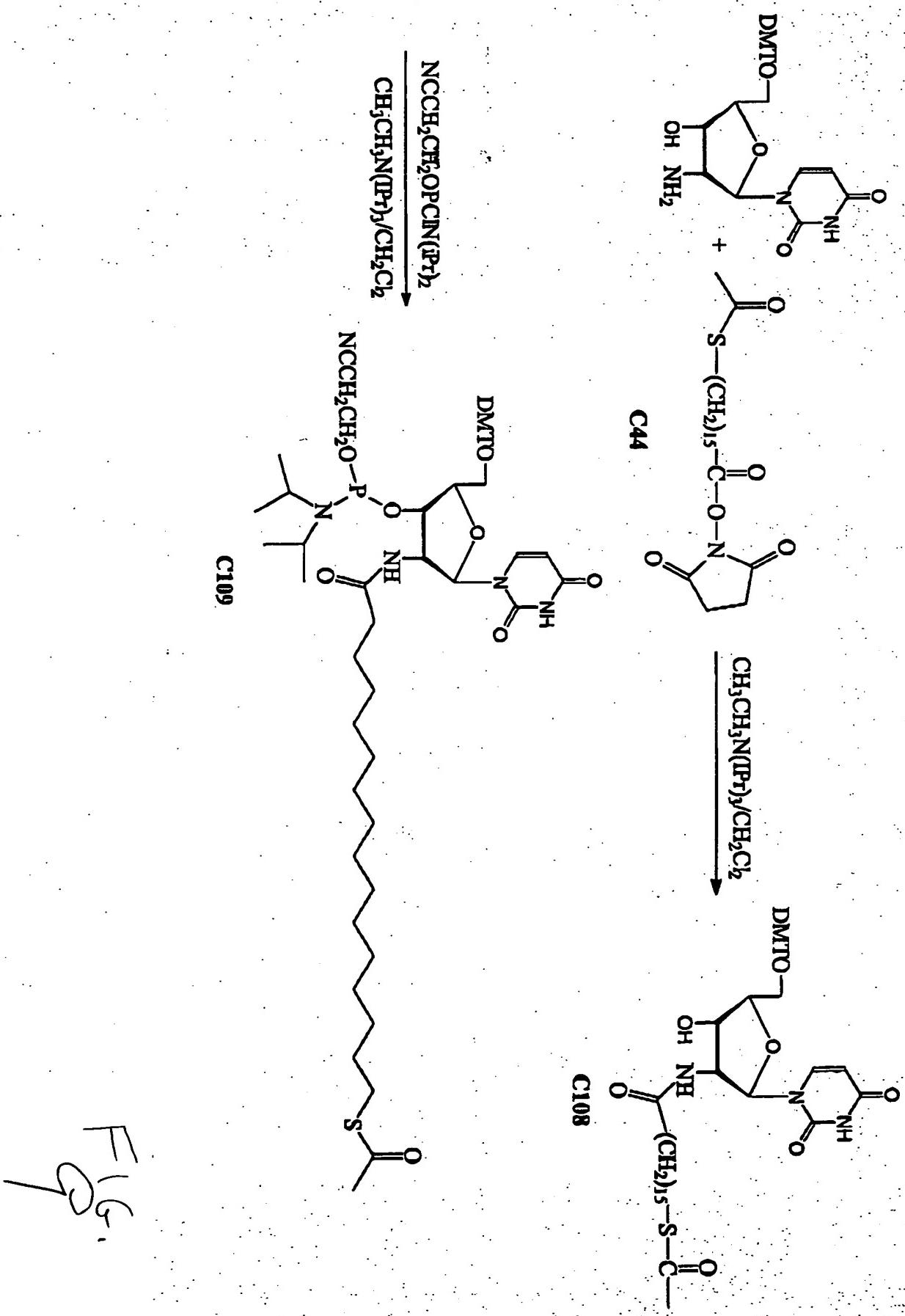
Any Position Attachment

8A

8B

FIG.  
8

## Synthesis Scheme of C109



**Synthesis Scheme f Ethylene Glycol Terminated Wires**

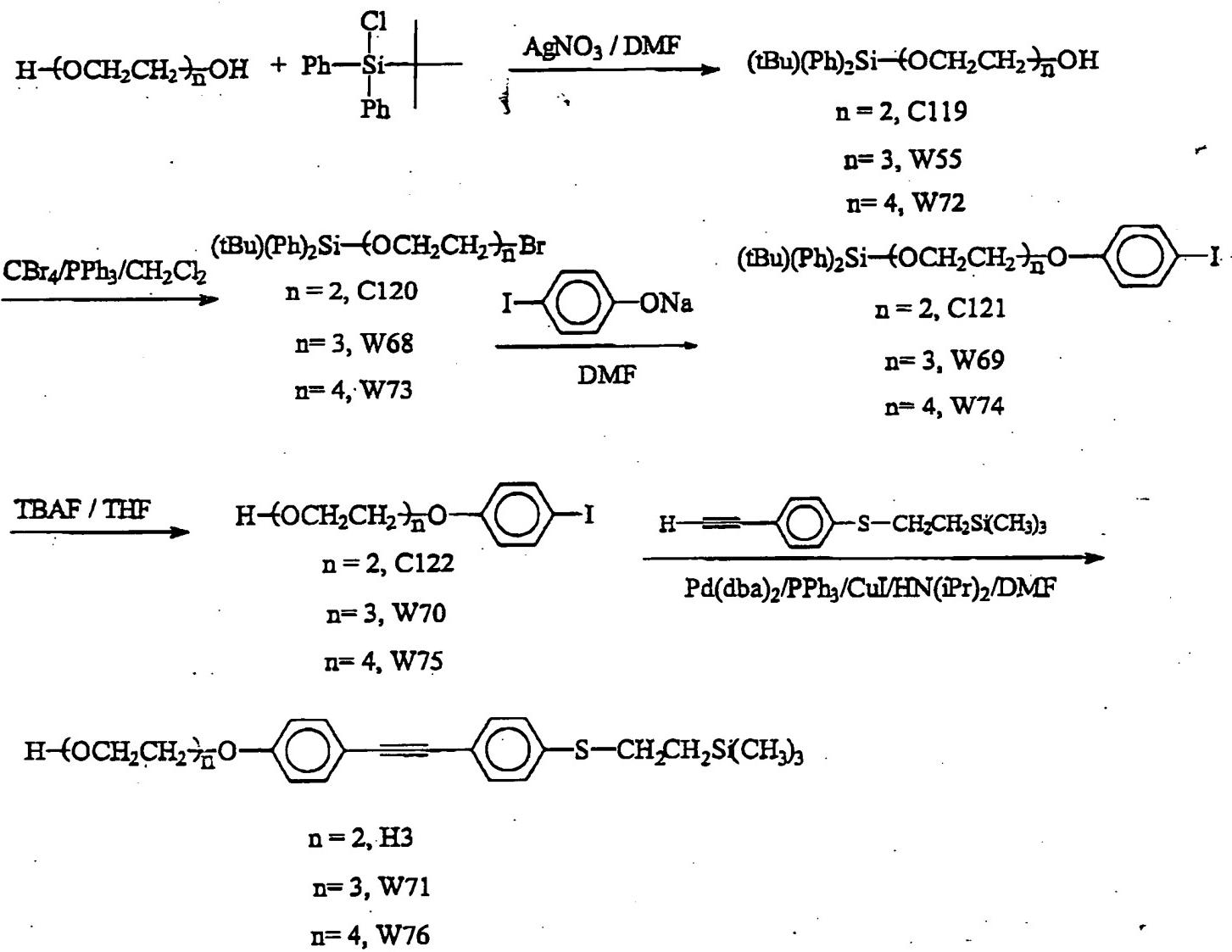


Fig.

10

## Synthesis Scheme of Branched Adenosine

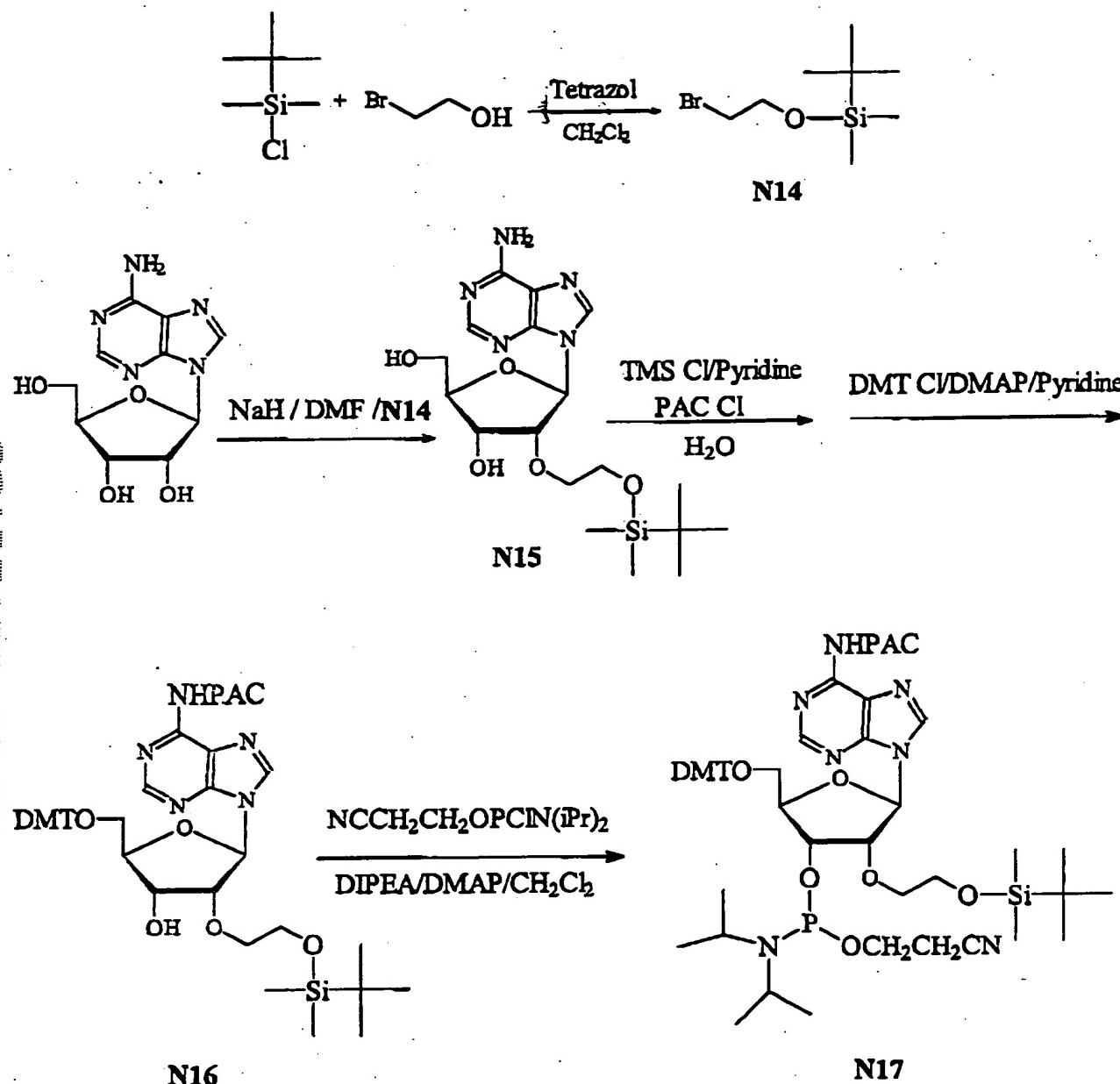


FIG.  
IIA

## Synthesis Scheme of W90

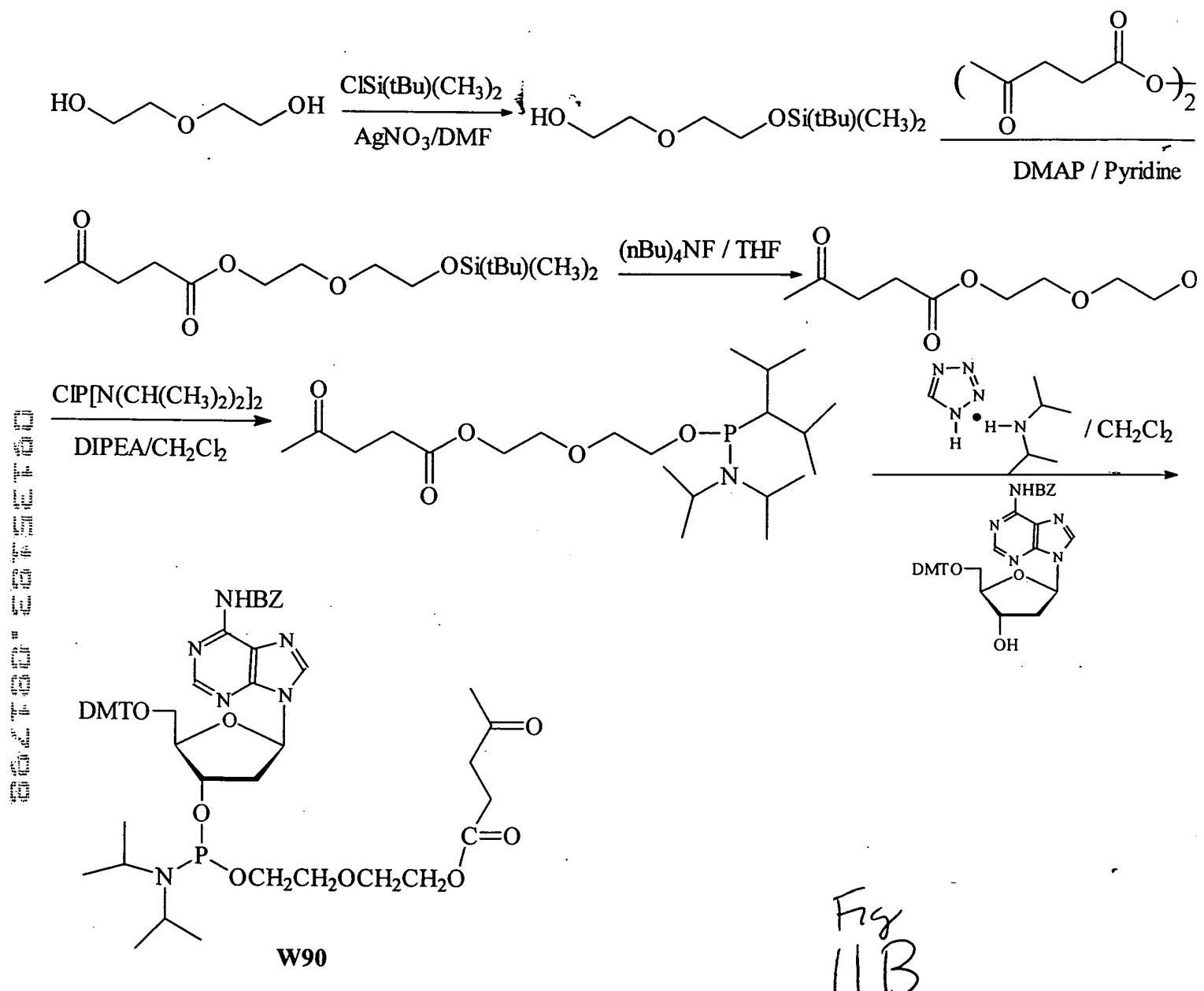


Fig  
11B

## Synthesis Scheme of N38

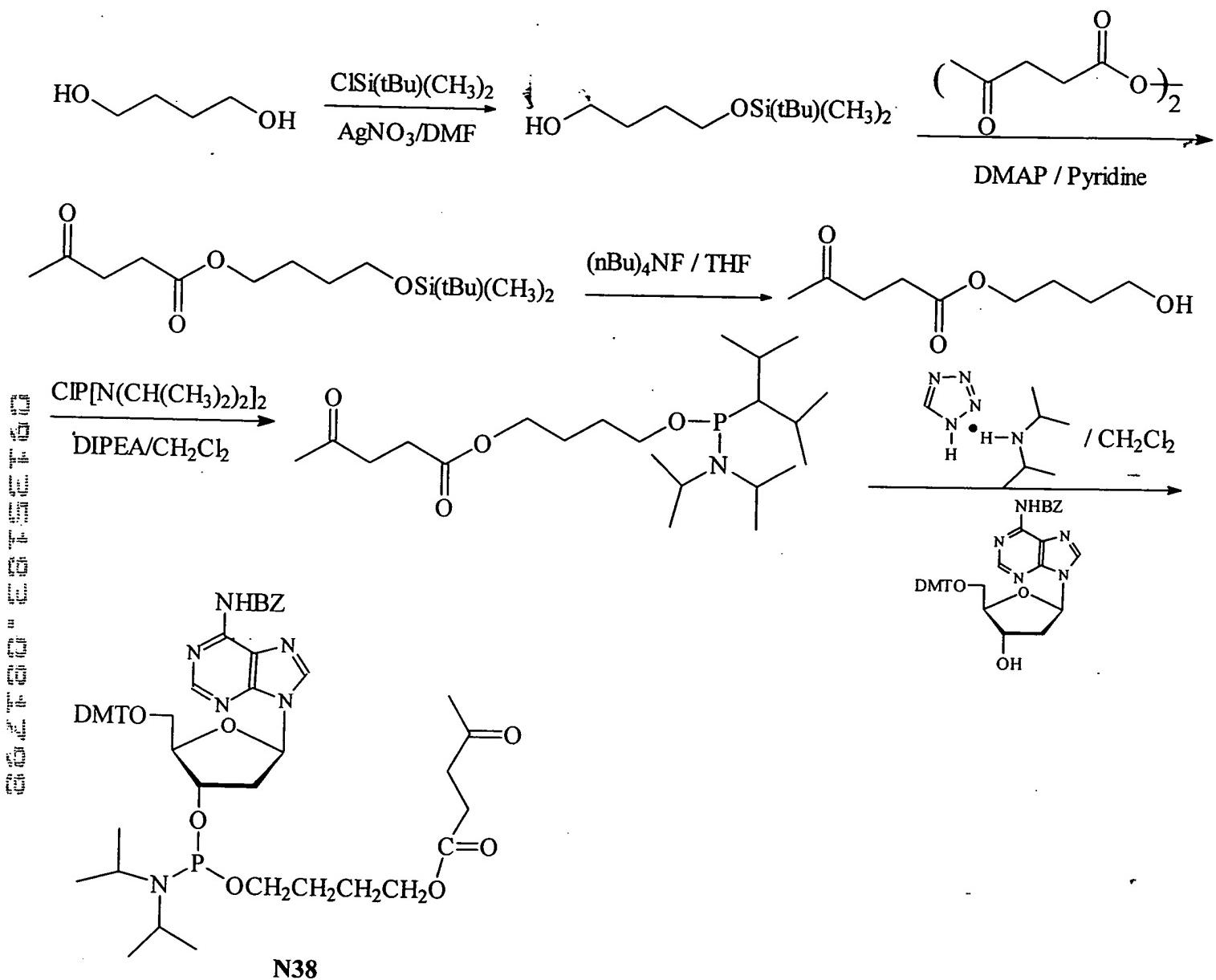
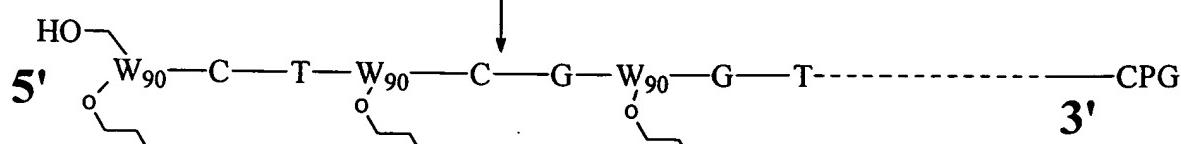


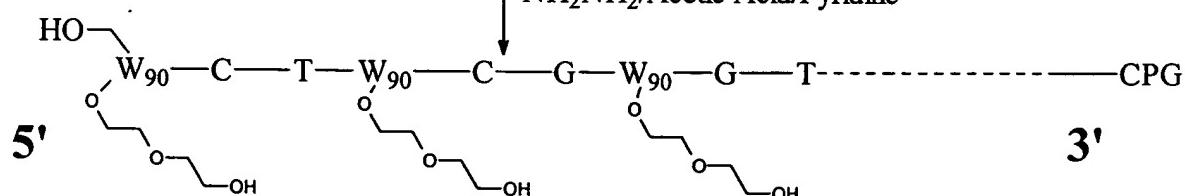
Fig  
11C

**Procedure of Incorporating Multiple Metal Complexes into DNA**

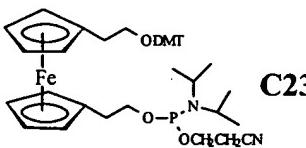
**Standard DNA Synthesis Using W90**



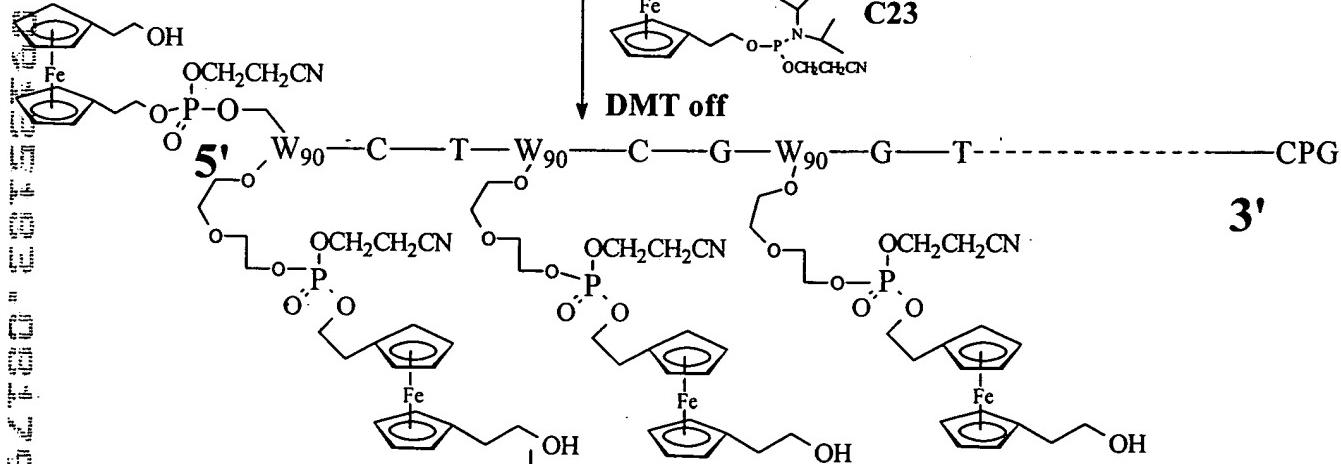
$\text{NH}_2\text{NH}_2/\text{Acetic Acid/Pyridine}$



**Coupling to**



**DMT off**



This process can be repeated until the desired # of Ferrocene is obtained, and then hydroxy groups on ferrocene are capped using the left phosphoramidite in order to increase the solubility of Ferrocene in water

**DMT off / Cleavage and Deprotection**

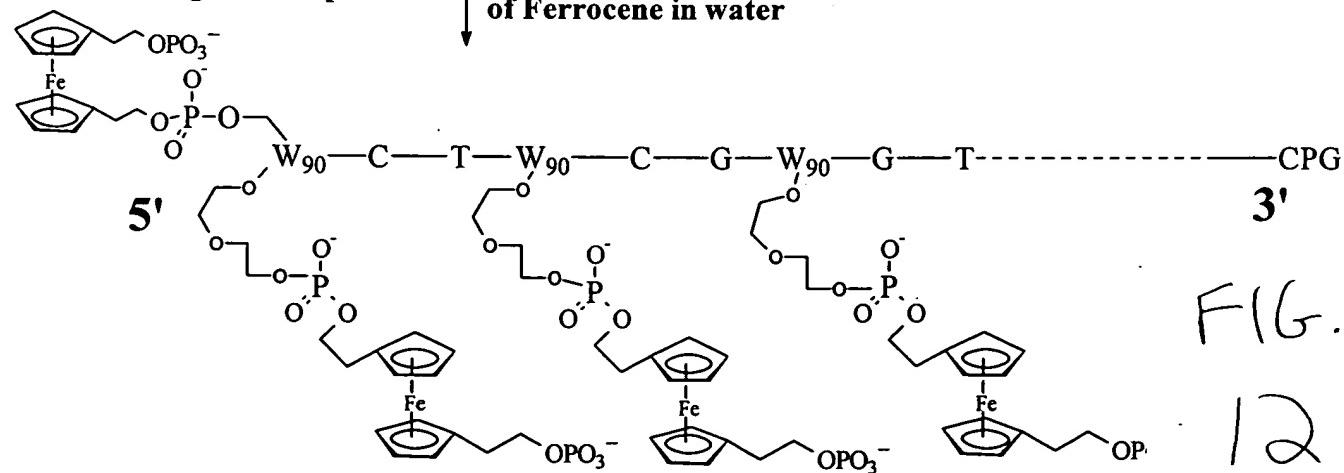
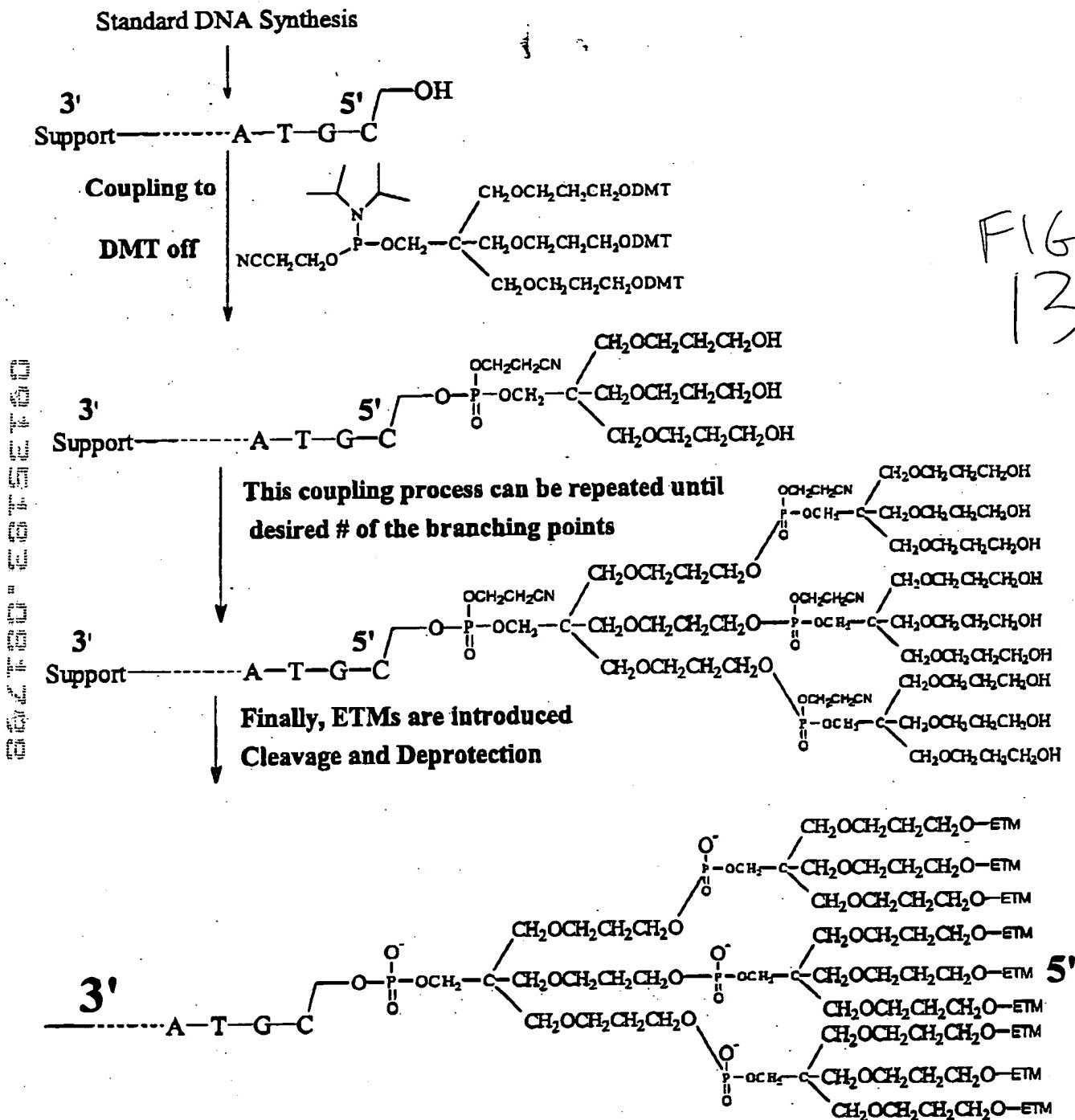


FIG.

12

### Scheme f Inc rp rating Multiple ETMs Using Branching Ph sphoramide



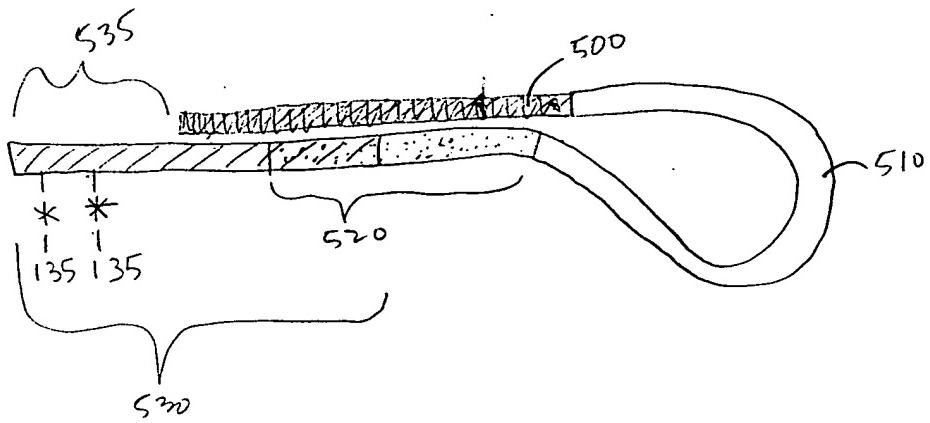
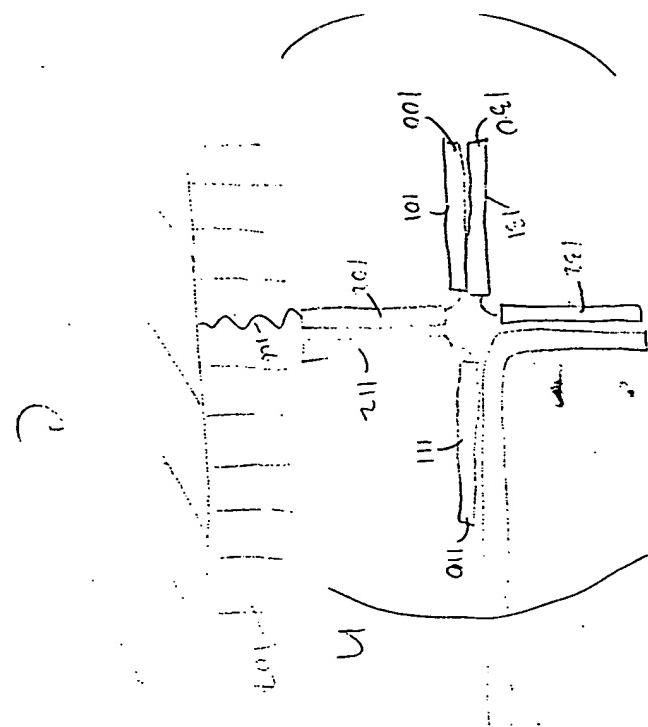
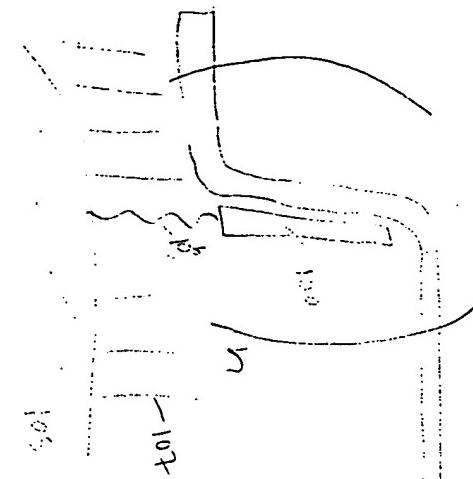


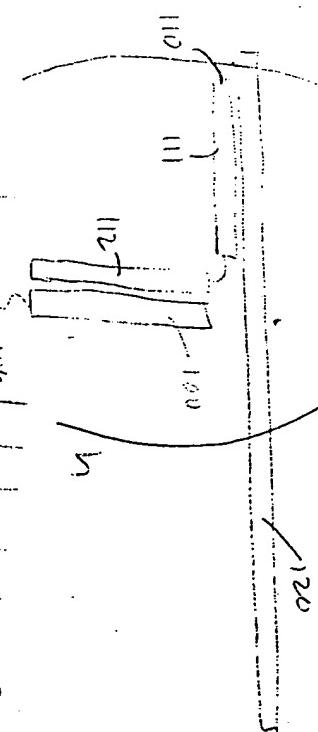
FIG. 14



4



170



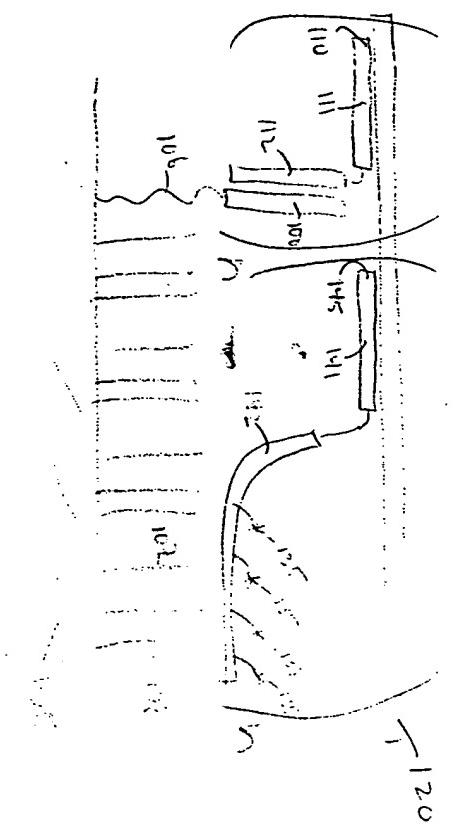
2

Fig.

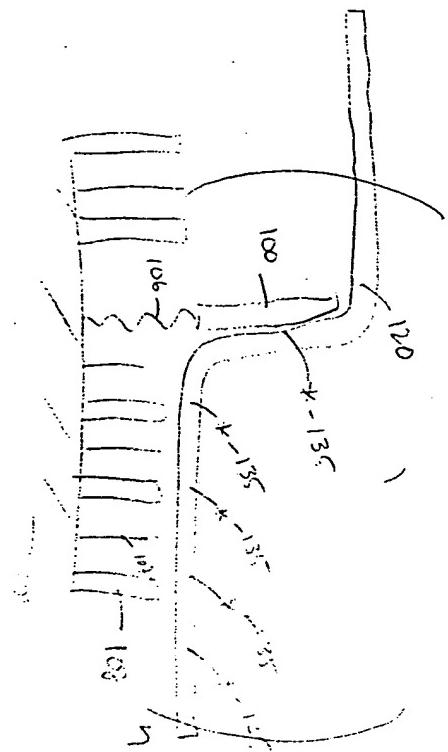
0 9 8 7 6 5 4 3 2 1

F1(5)6

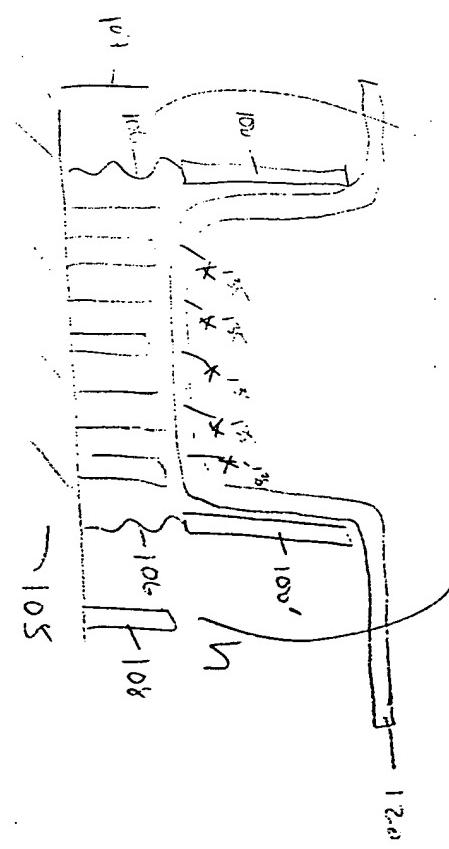
C



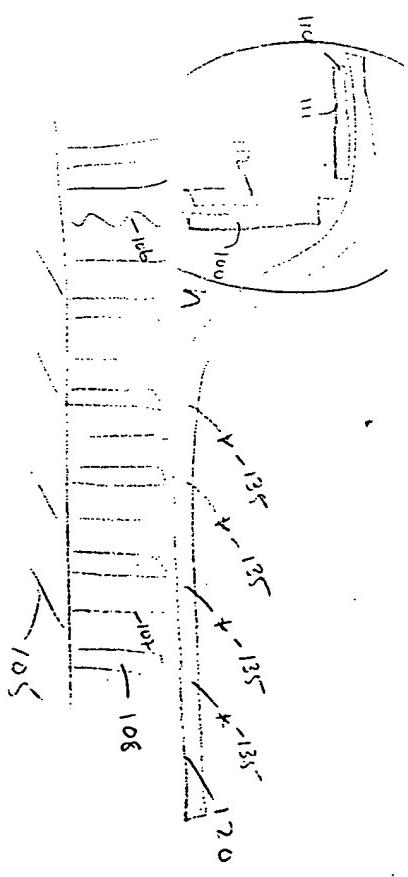
A



D



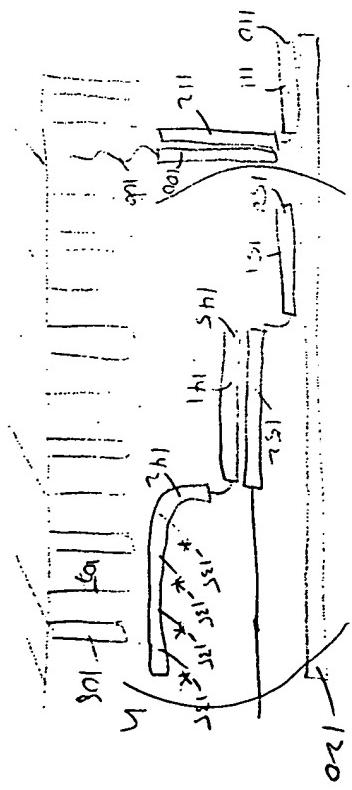
B



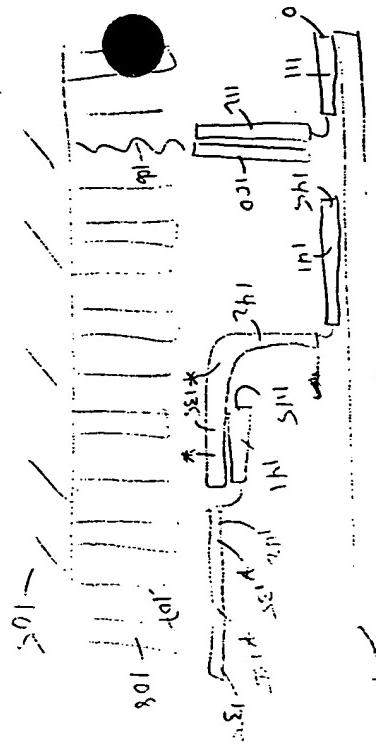
CONT'D

FIG.

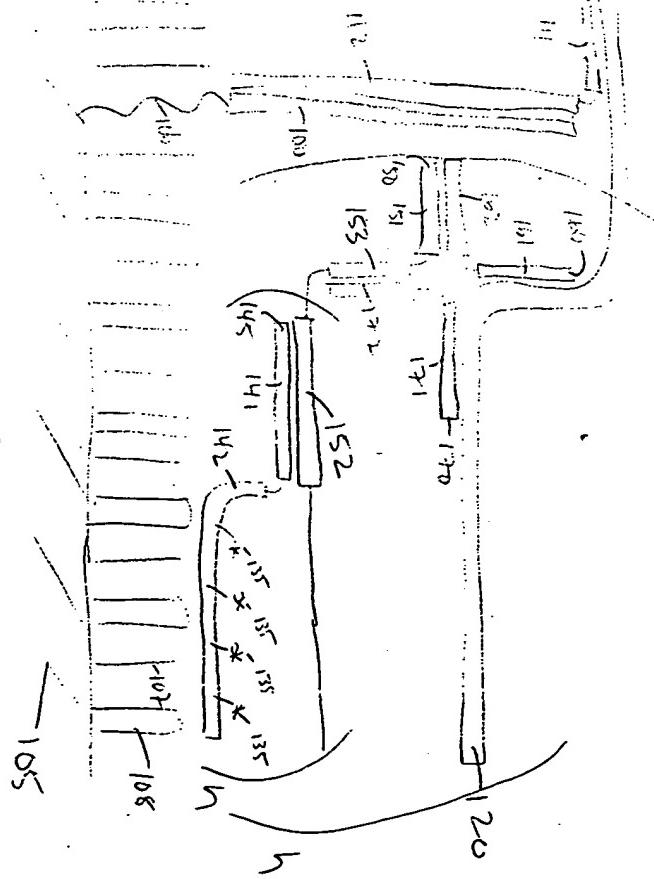
E



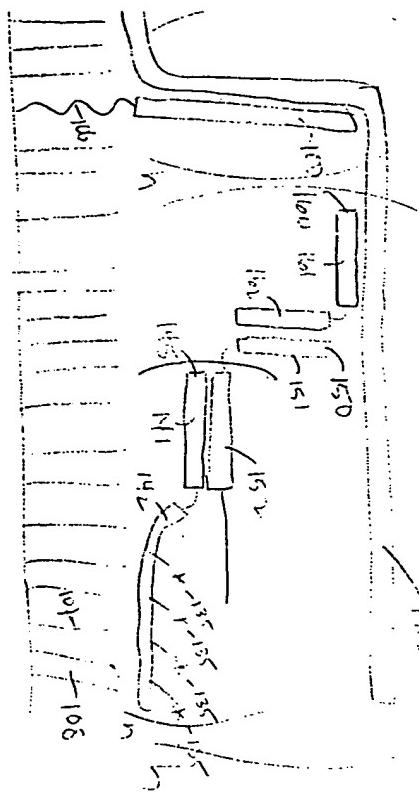
H



G



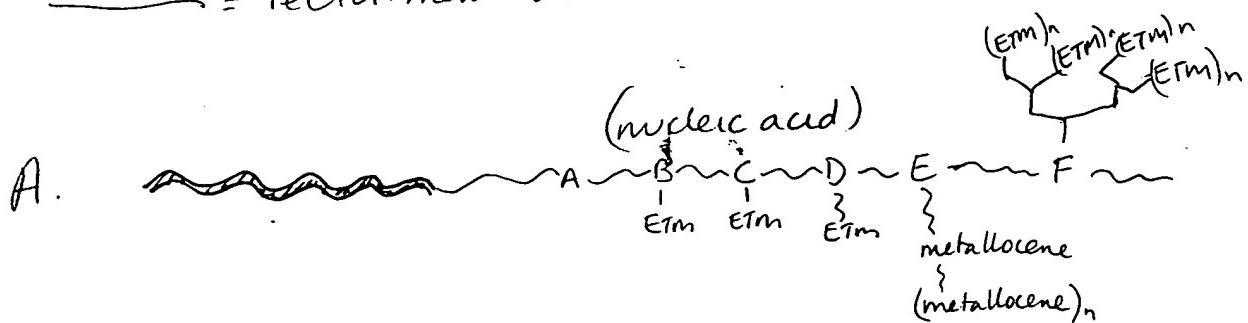
F



# Label probes

       = first <sup>hybridizable</sup> portion of label probe

       = recruitment linker



A = nucleoside replacement

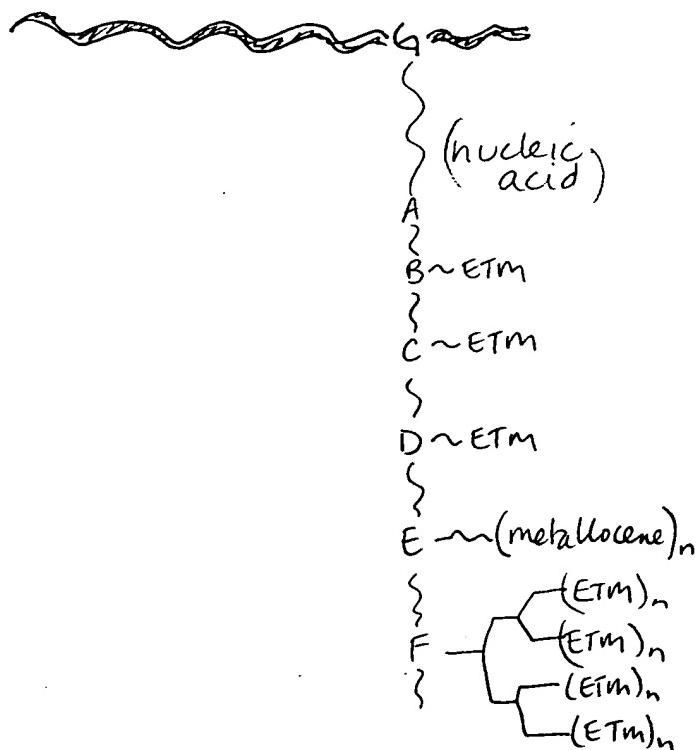
B = attachment to a base

C = attachment to a ribose

D = attachment to a phosphate

E = metallocene polymer, attached to a ribose, phosphate, or base

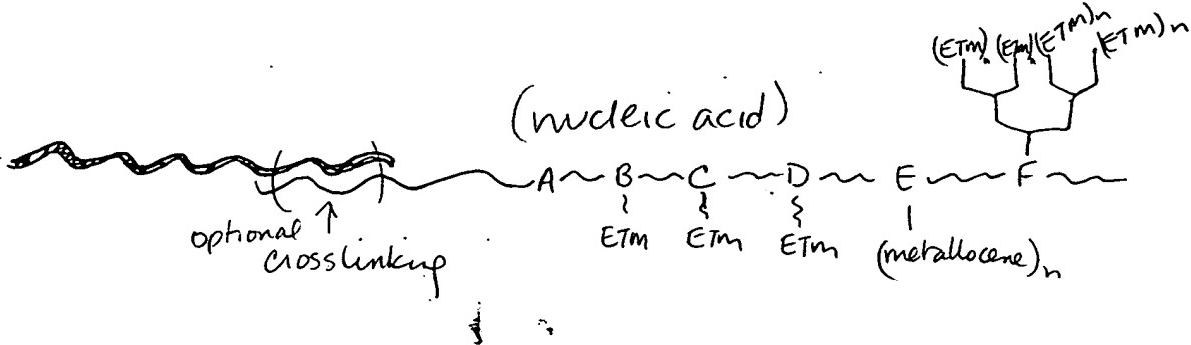
F = dendrimer structure, attached via a ribose, phosphate or base



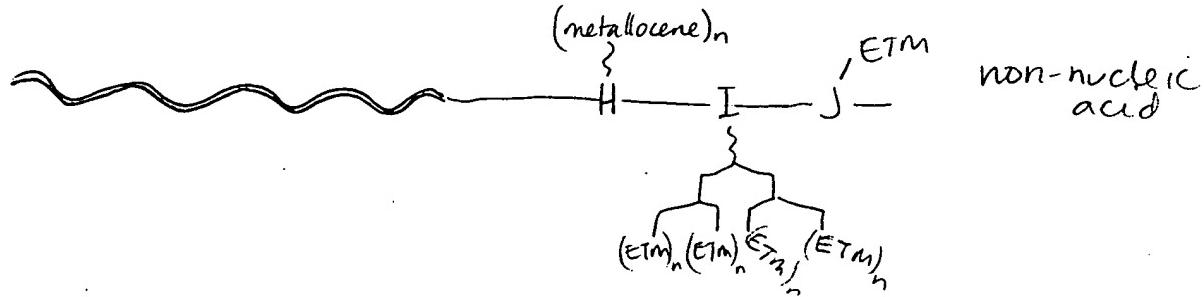
G = attachment via a "branching structure", through ribose, phosphate or base

FIG.  
17

C.



D.



H = attachment of metallocene polymers

I = attachment via dendrimer structure

J = attachment using standard linkers

E.

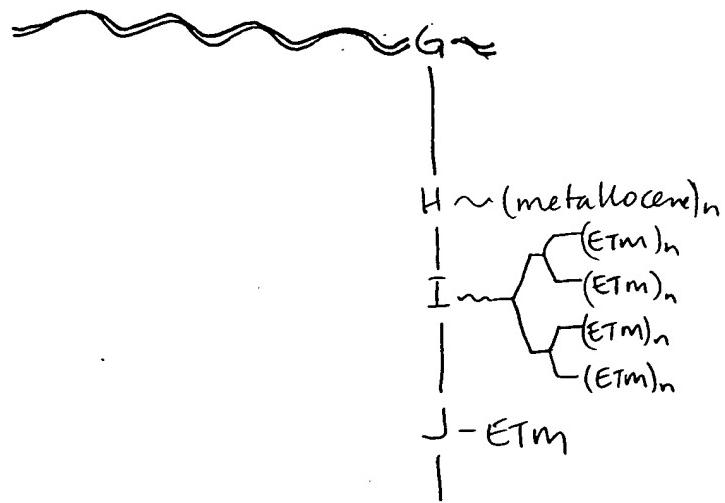


FIG.  
17  
cont

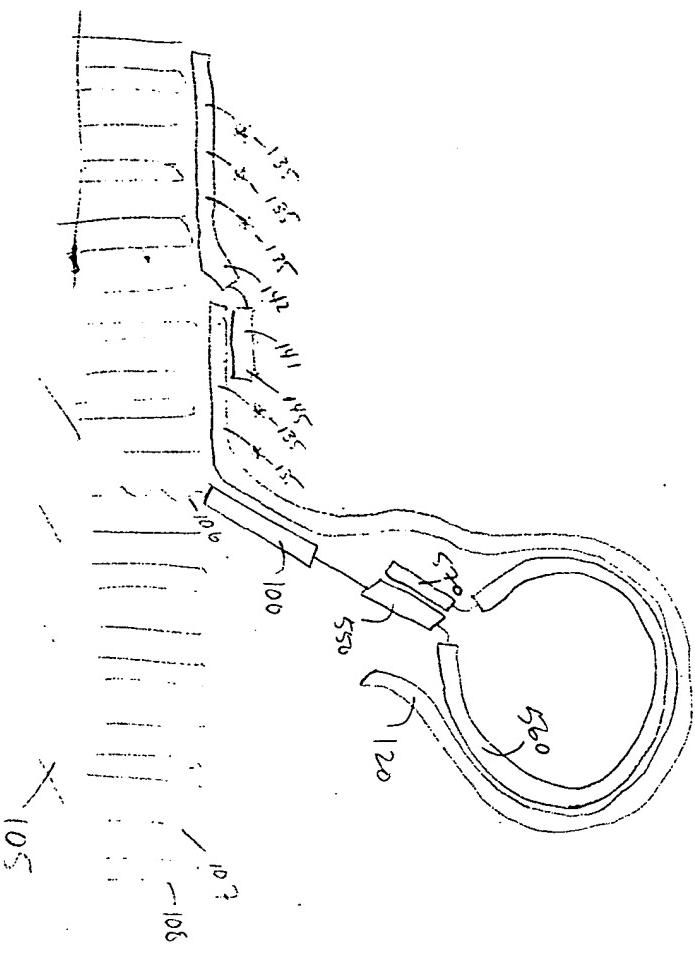


Fig.  
18.

D179  
5' - A(C15)CCTGGTCTTGACATCCACGGAAGGC<sup>G</sup>TGGAAATACGTATT<sup>C</sup>GT<sup>T</sup>GCCTA - 3'

D309 (Dendrimer)  
5' - (W38)(Branching)(Branching)CATGGTTAACGTCAATTGCTGCGGTTATTAA - 3'

D295  
5' - (N6)G(N6)CT(N6)C(N6)G(N6)C(N6)CCCATGGTTAGACTGAATTGCTGCGGTTATTAA - 3'

D297  
5' - (N6)G(N6)CT(N6)C(N6)G(N6)C(N6)TATGCTCTTATGGTGTGGAAATCTACTGG - 3'

D298  
5' - (N6)G(N6)CT(N6)C(N6)G(N6)C(N6)ATGGTGTGGAAATCTACTGG - 3'

D296  
5' - (N6)G(N6)CT(N6)C(N6)G(N6)C(N6)TGACTGAATTGCTGCGGTTATTAA - 3'

D112  
5' - CTTCCGTGGATGTCAAGACCAGGAU - 4 unit wire (C11) - 3'

D94  
5' - ACCATGGACACAGAU - 4 unit wire (C11) - 3'

D109  
5' - CTGCGGTTATTAAACU - 4 unit wire (C11) - 3'

2Tar.  
5' - TAG GCA CGA ATA CGT ATT TCC ACG ATA AAT ATA ATT AAT AAC CGC AGC AAT TGA CGT ATA AAG CTA TCC CAG TAG ATT TCC ACA GC - 3'

D349  
5' - A(C15)C (C15)GT GTC CAT GGT AGT AGC TTA TCG TGG AAA TAC GTA TTC GTG CCT A - 3'

D382  
5' - (Y63)G(Y63) CT(Y63) C(Y63)G (Y63)C(Y63) CCC ATG GTT AGA CTG AAT TGC TGC GGT TAT TAA - 3'

D383  
5' - (Y63)G(Y63) CT(Y63) C(Y63)G (Y63)C(Y63) CCC ATG GTT AGA CTG GCT GTG GAA ATC TAC TGG - 3'

D468  
5' - (N6)G(N6) CT(N6) C(N6)G (N6)C(N6) (glen)(glen)(glen) CTT TAC TCC CTT CCT CCC CGC TGA AAG TAC - 3'

D449  
5' - CGG AGT TAG CCG GTG CTT CTT CTG CGG G(C131)(C131) (C131)(C131)(N6) G(N6)C T(N6)C (N6)G(N6) C(N6)T - 3'

D417  
5' - CTT TAC TCC CTT CCT CCC CGC TGA AAG TAC TTT ACA ACC C - 3'

F16  
F17

EUI

5' - ATC CTG GTC TTG ACA TCC ACG GAA GAT GTC CCT ACA GTC TCC ATC AGG CAG TTT  
CCC AGA CA - 3'

MT1

5' - TCT ACA TGC CGT ACA TAC GGA ACG TAC GGA GCA TCC TGG TCT TGA CAT CCA CGG  
AAG - 3'

D358

5' - (N6)G(N6) CT(N6) C(N6)G (N6)C(N6) CCG TAT GTA CGG CAT GTA GA - 3'

D334

5' - GCT ACT ACC ATG GAC ACA GAU - 4 unit wire (C11) - 3'

D335

5' - ACA GAC ATC AGA GTA ATC (N6)GC C(N6)G TC(N6) TGG (N6)T - 3'

LP280

5' - GAT TAC TCT GAT GTC TGT CCA TCT GTG TCC ATG GTA GTA GC - 3'

LN280

5' - GAT TAC TCT GAT GTC TGT CCT AGT ACG AGT CAG TCT CTC CA - 3'

NC112

5' - TCT ACA TGC CGT ACA TAC GGA ACG TAC GGA GCG ATT CGA CTG ACA GTC GTA ACC  
TCA - 3'

D336

5' - (N6)G(N6) CT(N6) C(N6)G (N6)C(N6) GCG ACA ACT GTA CCA TCT GTG TCC ATG GT - 3'

D405

5' - (C23)(C23)(C23) (C23)(C23)(C23) (C23)(C23)(C23) (C23)AT CTG TGT CCA TGG T - 3'

D429

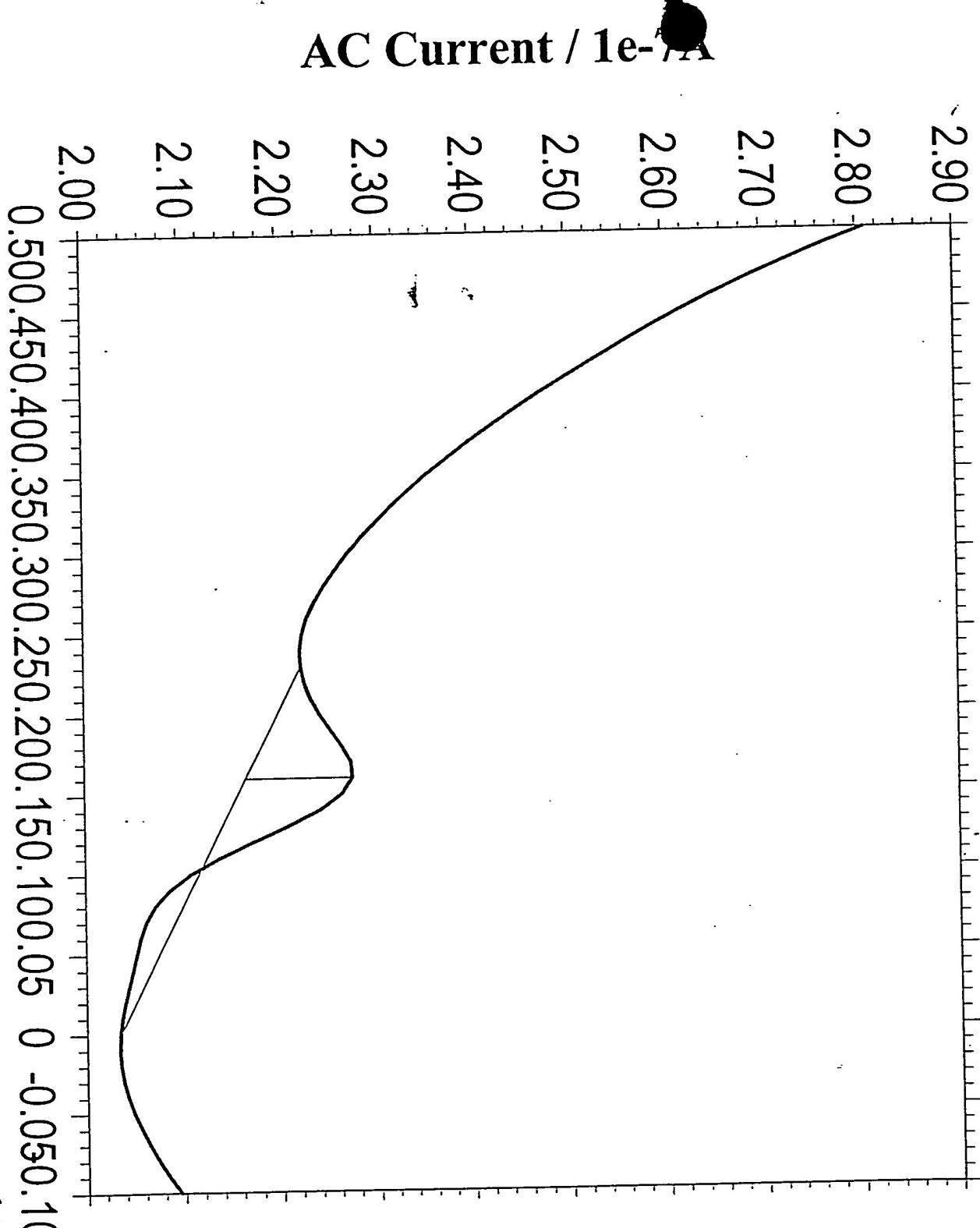
5' - (N6)G(N6) CT(N6) C(N6)G (N6)C(N6) (C131)AT CTG TGT CCA TGG TAG TAG C - 3'

Fig.  
19 cont

**Electrode # 55, d179+2tar+309+10%ACN**

Mar. 19, 1998 17:18:47  
Tech: ACV  
File: a292\_023

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7  
 $E_p = 0.160V$   
 $i_p = 1.092e-8A$   
 $\Delta p = 7.563e-10VA$



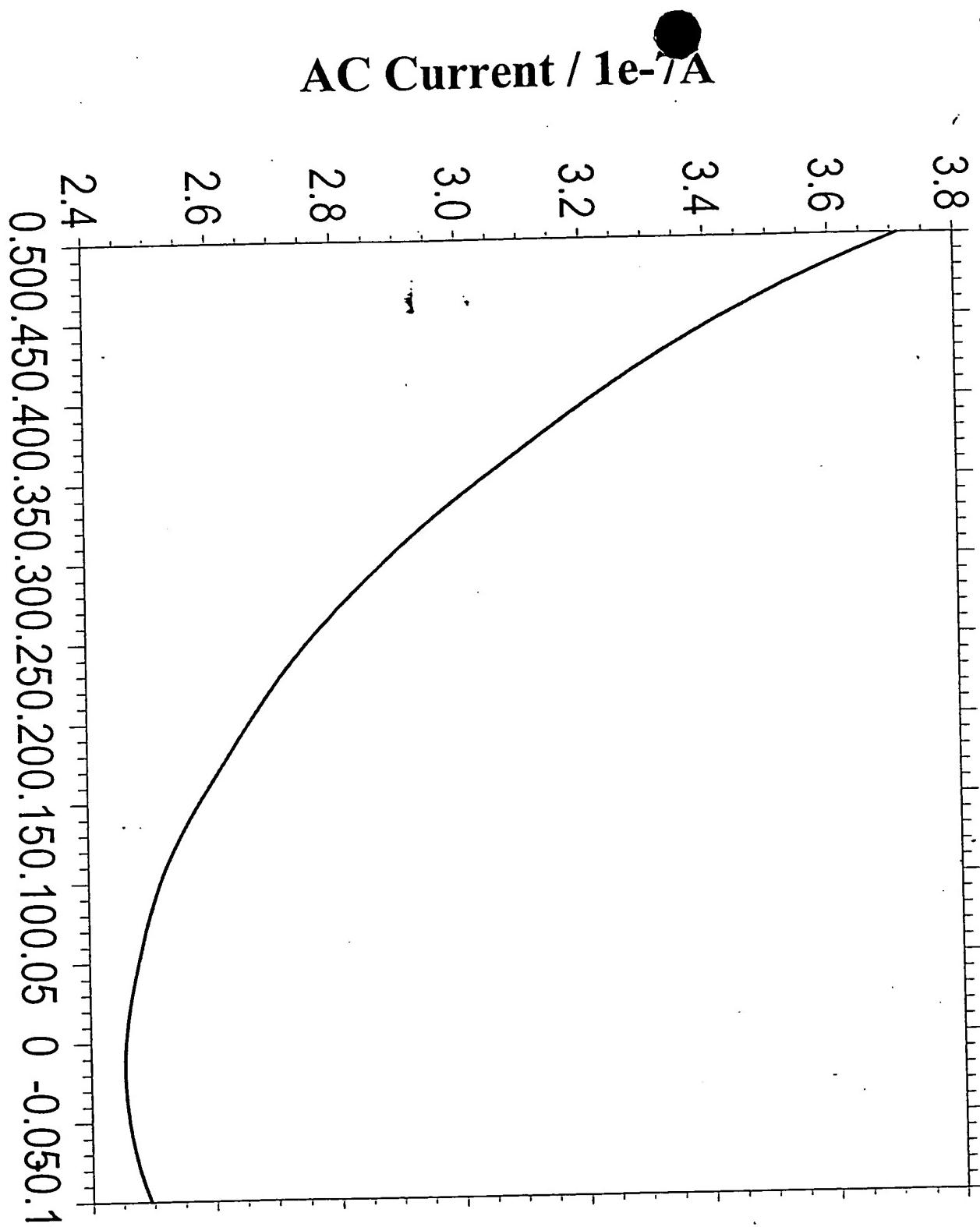
**CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.**

Electrode # 44, d1 / 9+30y+10%AUW

Mar. 19, 1998 17:00:25

Tech: ACV  
File: a292\_019

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7



CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

electrode #25

May 14, 1998 16:37:13  
Tech: ACV  
File: a358\_009

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

$E_p = 0.190V$   
 $i_p = 2.046e-7A$   
 $A_p = 2.046e-8VA$

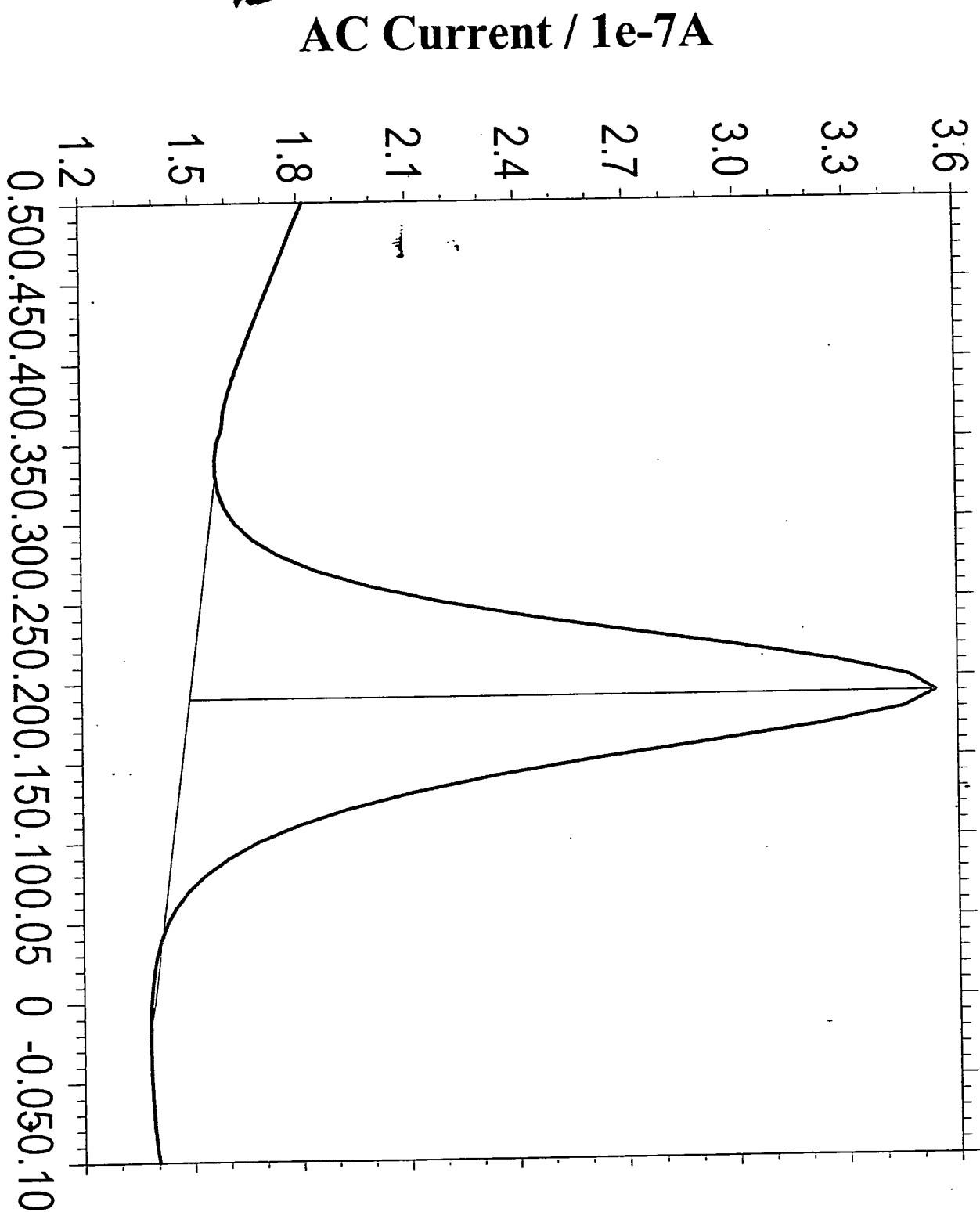


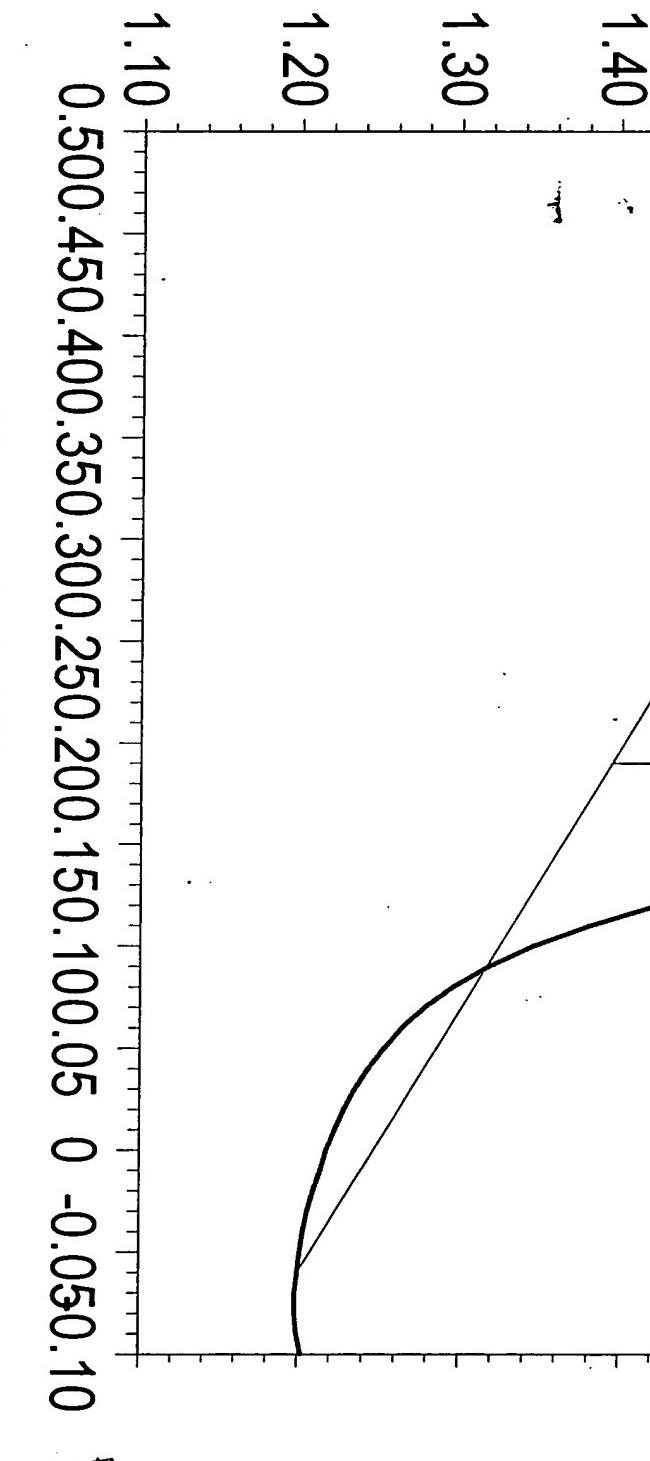
Fig.  
20 C

**electrode #37**

May 14, 1998 16:58:47  
Tech: ACV  
File: a358\_013

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7  
 $E_p = 0.190V$   
 $i_p = 3.552e-8A$   
 $A_p = 3.568e-9VA$

**AC Current /  $1e^{-7}A$**

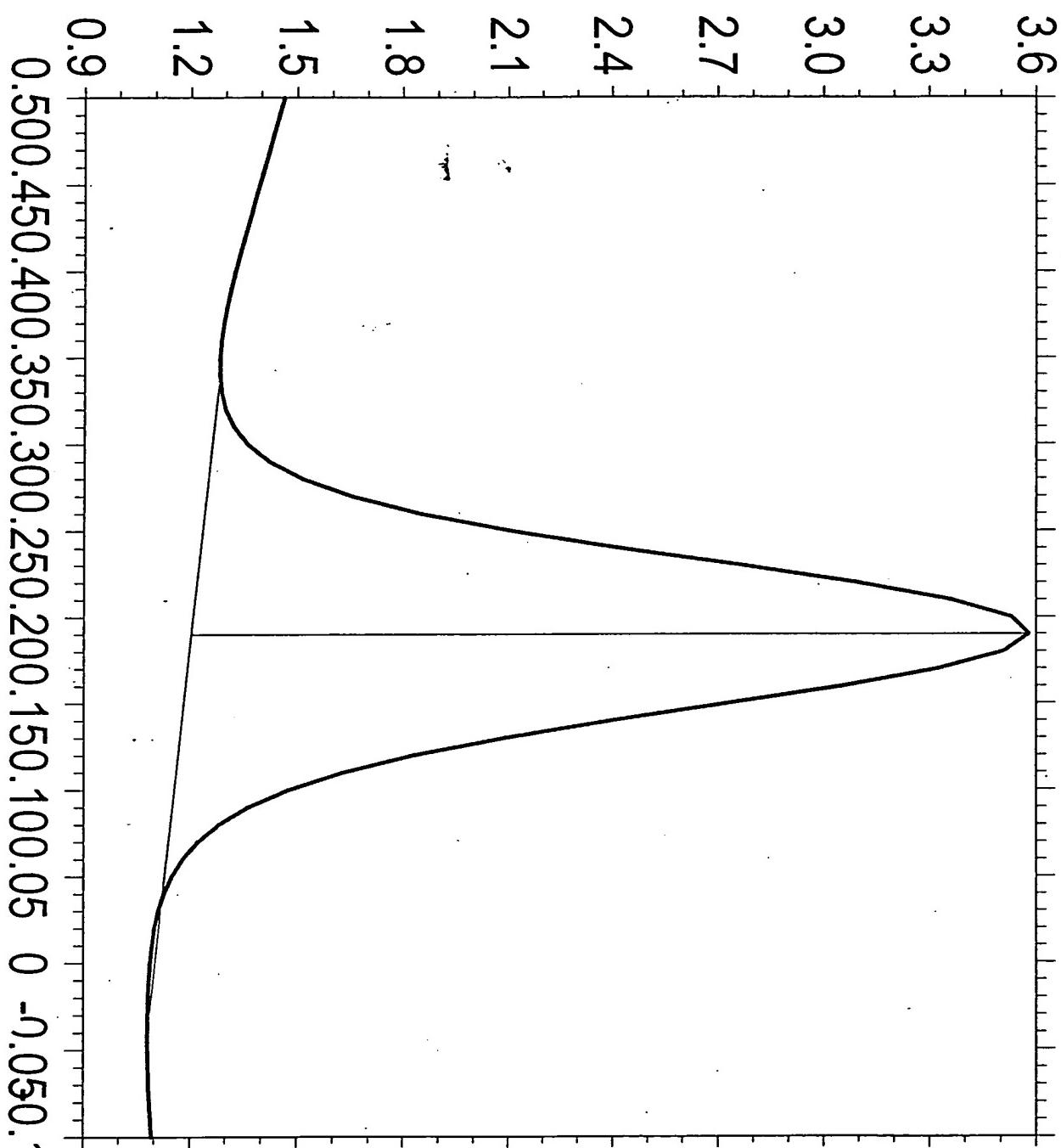


**Potential / V vs Ag/AgCl**

f16.  
20 D

# Electrode 6

AC Current /  $1e-7A$



May 14, 1998 15:59:14  
Tech: ACV  
File: z039\_002

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7  
 $E_p = 0.190V$   
 $i_p = 2.376e-7A$   
 $A_p = 2.594e-8VA$

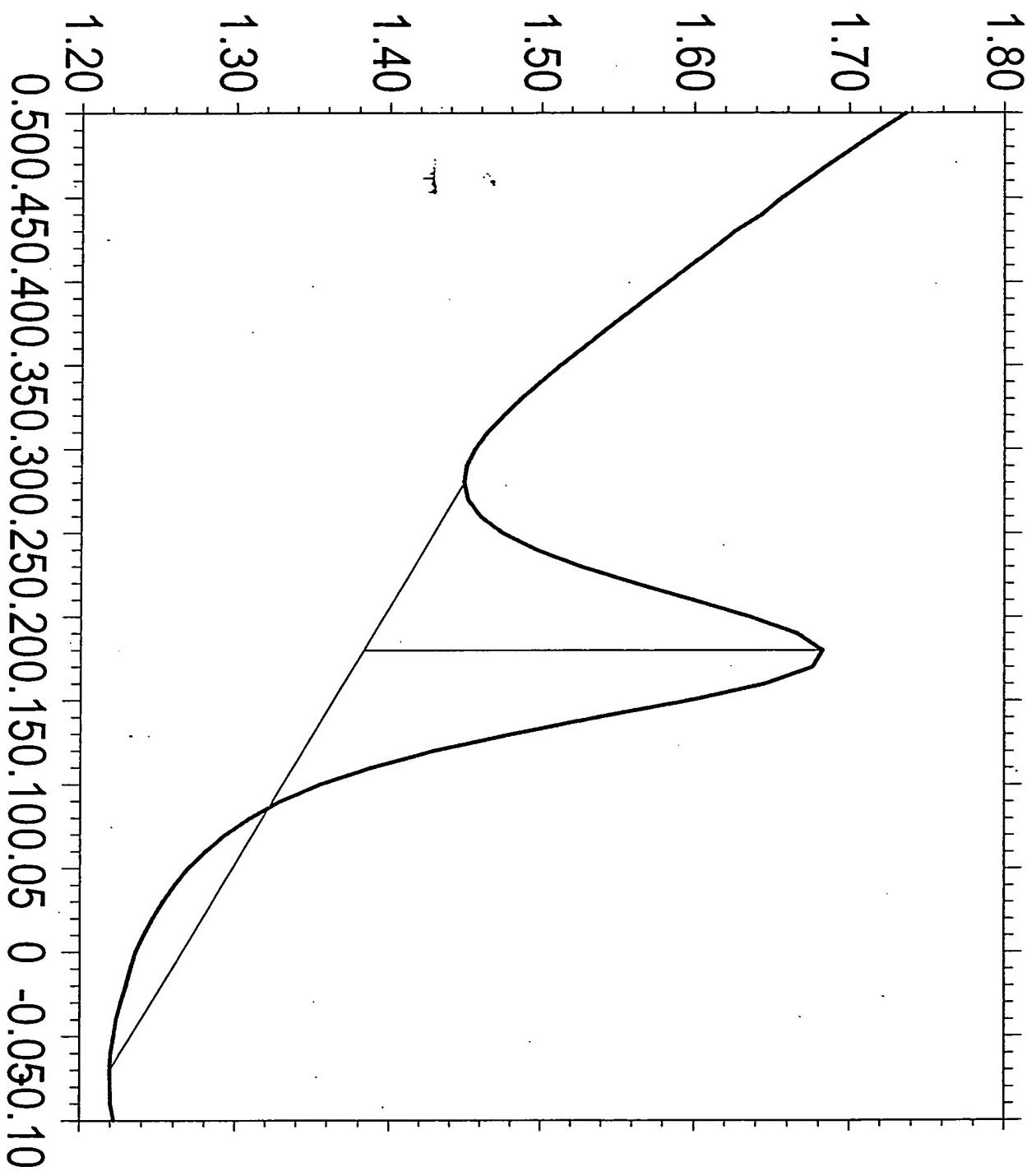
CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

Potential / V  
F16.20

electrode #40

14

AC Current /  $1e-7A$



CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

Potential / V vs Ag/AgCl

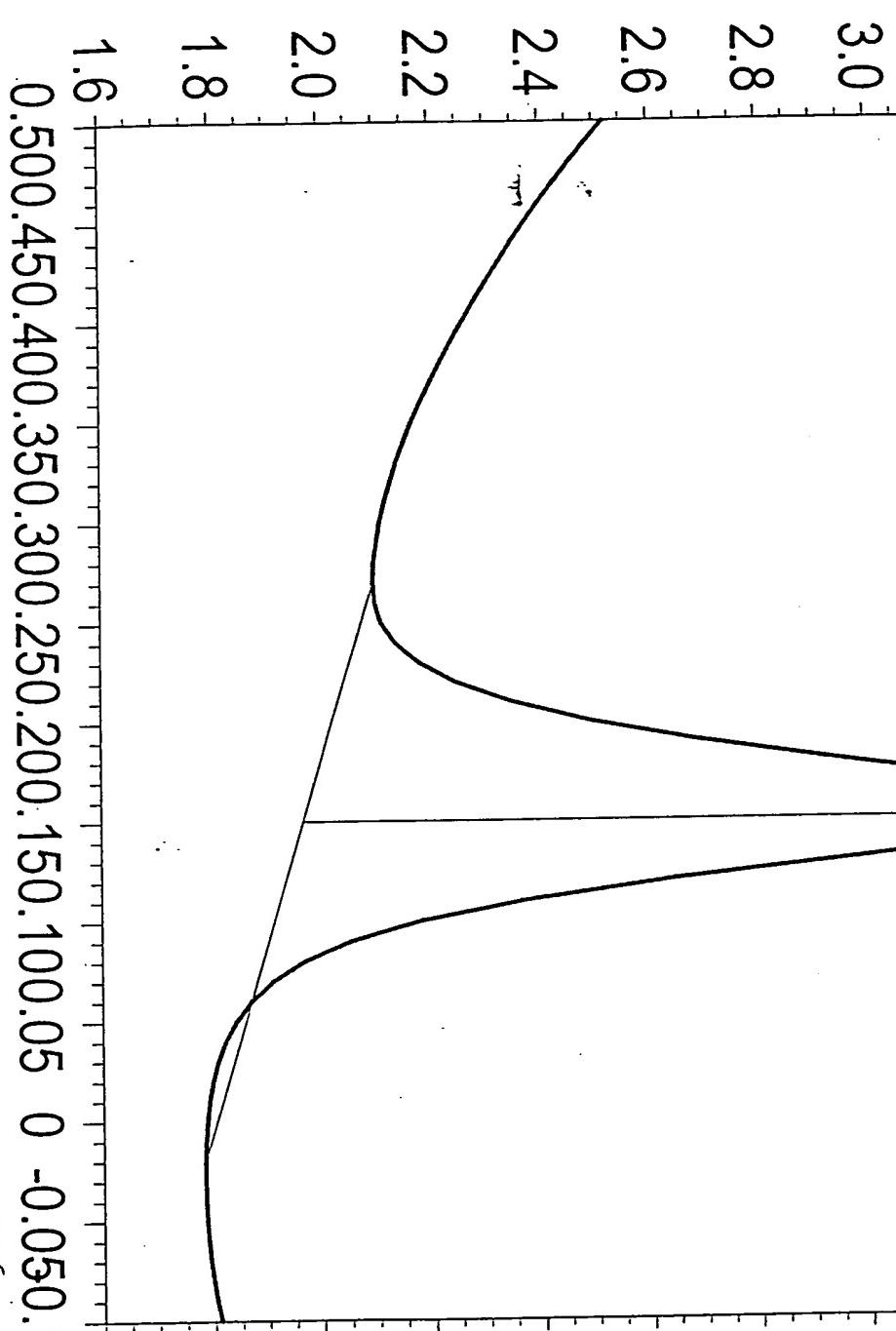
F16  
F20

# Electrode # 16

Mar. 19, 1998 16:00:02  
Tech: ACV  
File: v368\_028

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

$E_p = 0.150V$   
 $i_p = 1.494e-7A$   
 $A_p = 1.100e-8VA$



CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

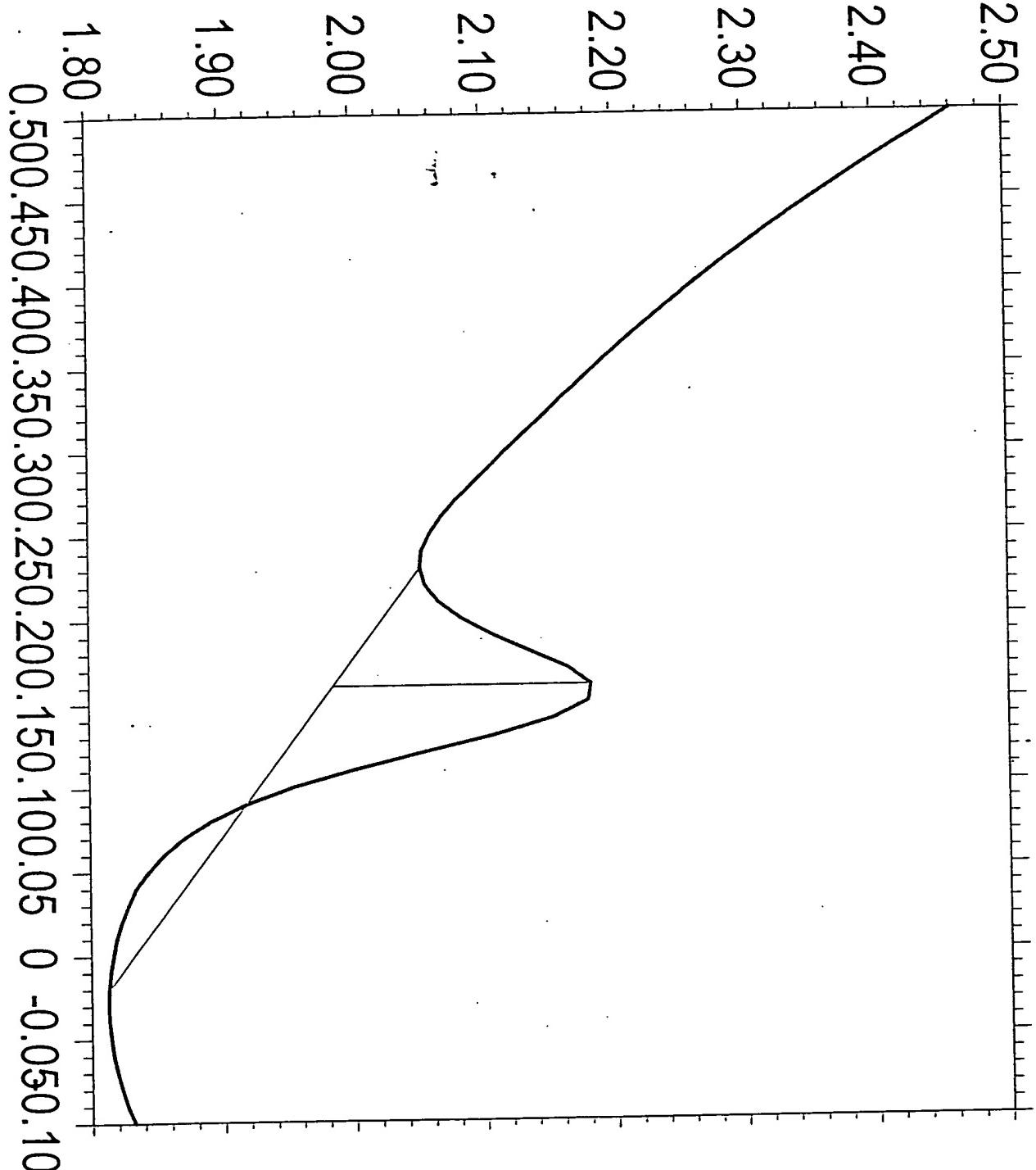
**Electrode # 18**

Mar. 19, 1998 16:17:15  
Tech: ACV  
File: v368\_032

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

$E_p = 0.160V$   
 $i_p = 1.967e-8A$   
 $A_p = 1.443e-9VA$

**AC Current /  $1e^{-7}A$**



Potential / V vs Ag/AgCl

F16.  
Q20 H

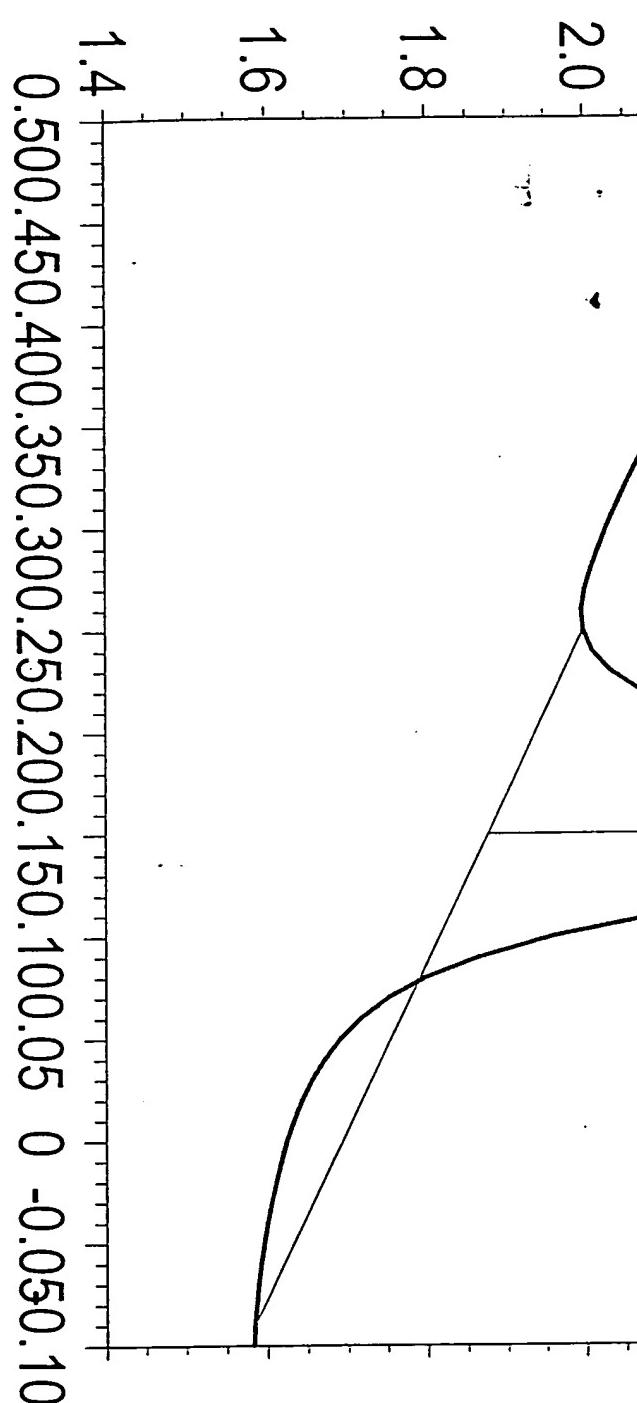
**CONFIDENTIAL PROPERTY OF**  
**Clinical Micro Sensors, Inc.**

**Electrode # 17**

Mar. 19, 1998 16:13:12  
Tech: ACV  
File: v368\_031

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7  
 $E_p = 0.150V$   
 $i_p = 8.031e-8A$   
 $A_p = 6.033e-9VA$

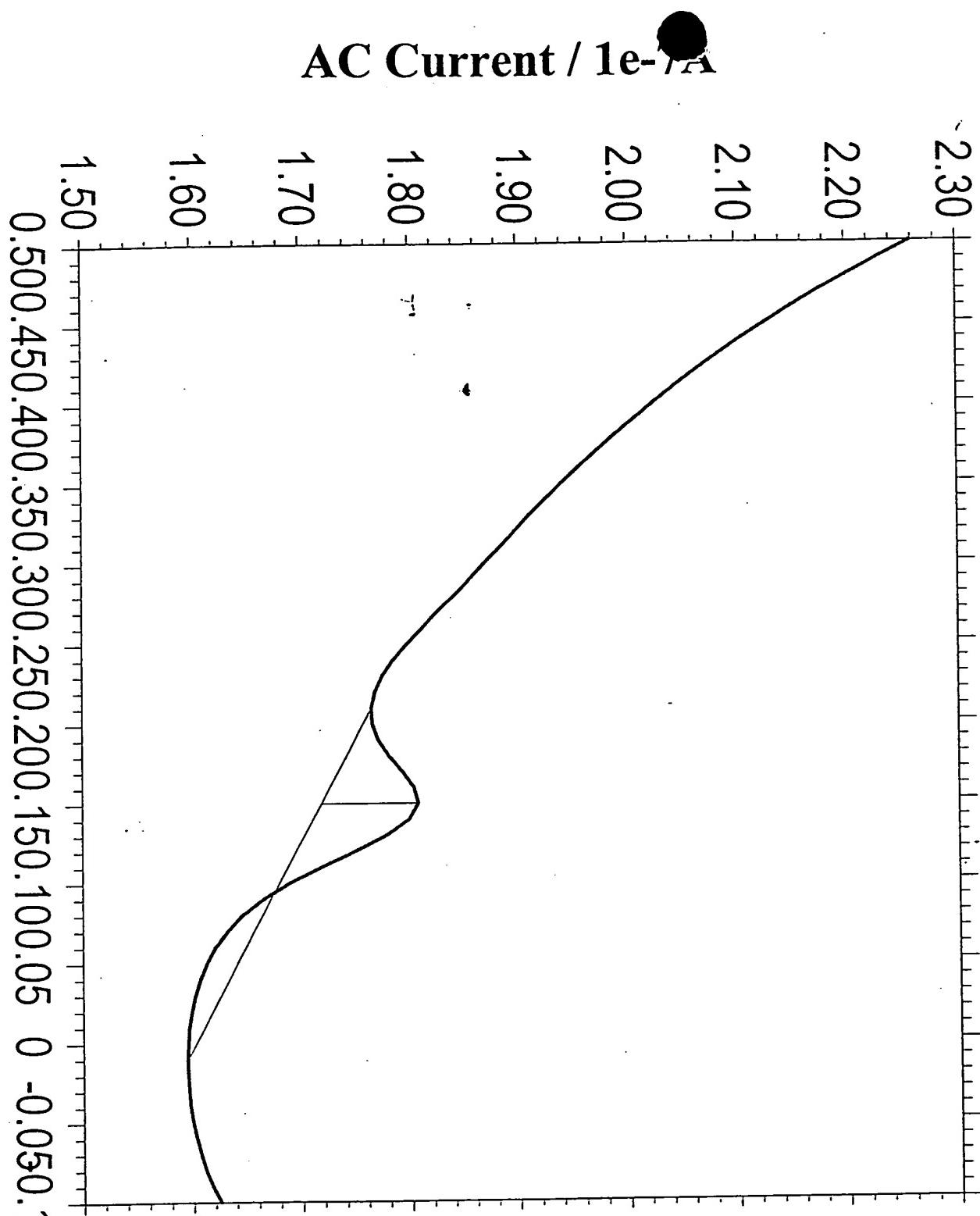
**AC Current / 1e-7A**



Potential / V vs Ag/AgCl  
0.500 0.450 0.400 0.350 0.300 0.250 0.200 0.150 0.100 0.050 0.000  
1.4 1.6 1.8 2.0 2.2 2.4 2.6

**CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.**

# Electrode # 13



Mar. 19, 1998 15:30:16  
Tech: ACV  
File: v368\_019

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

$E_p = 0.150\text{V}$   
 $i_p = 8.871\text{e-9A}$   
 $A_p = 5.512\text{e-10VA}$

CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

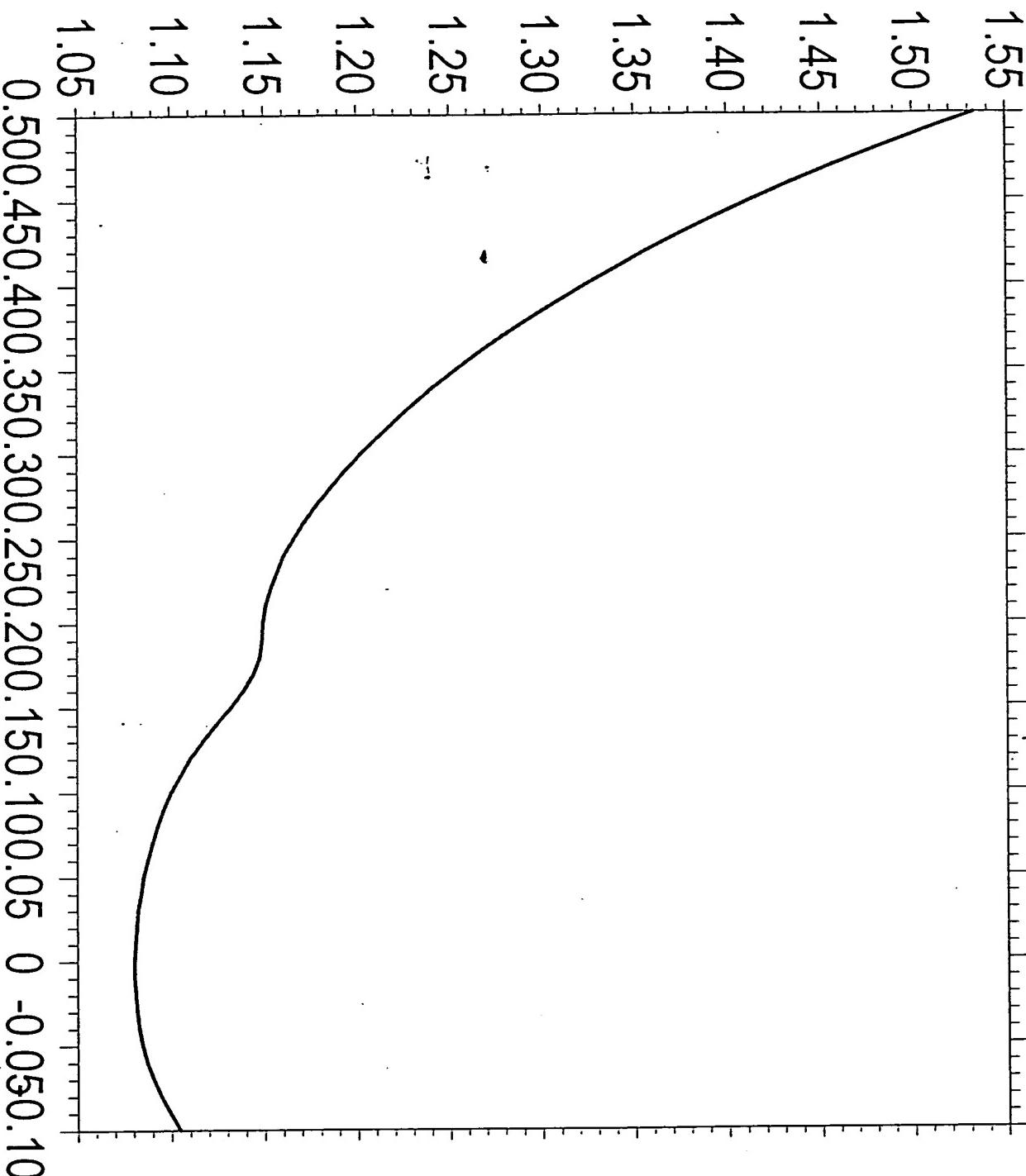
Electrode #22

13

AC Current /  $1e^{-7}A$

May 26, 1998 16:38:44  
Tech: ACV  
File: a371\_008

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) =  $2e^{-7}$



Potential / V vs Ag/AgCl  
F 16 K  
20

CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

# Electrode #15

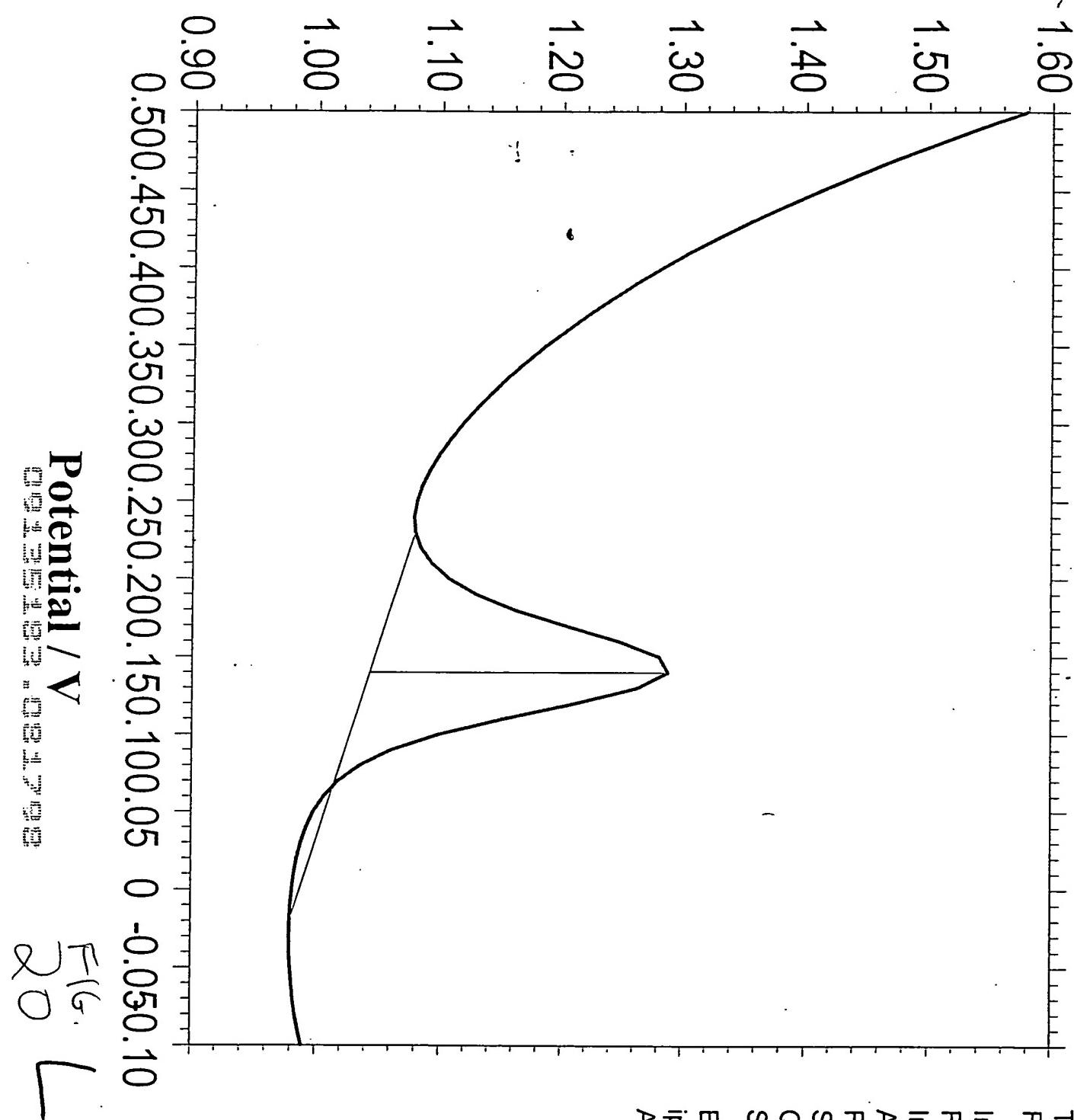
Apr. 6, 1998 13:58:20

Tech: ACV  
File: u059\_013

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 1e-6

$E_p = 0.140V$   
 $i_p = 2.449e-8A$   
 $A_p = 1.706e-9VA$

## AC Current / 1e-7A



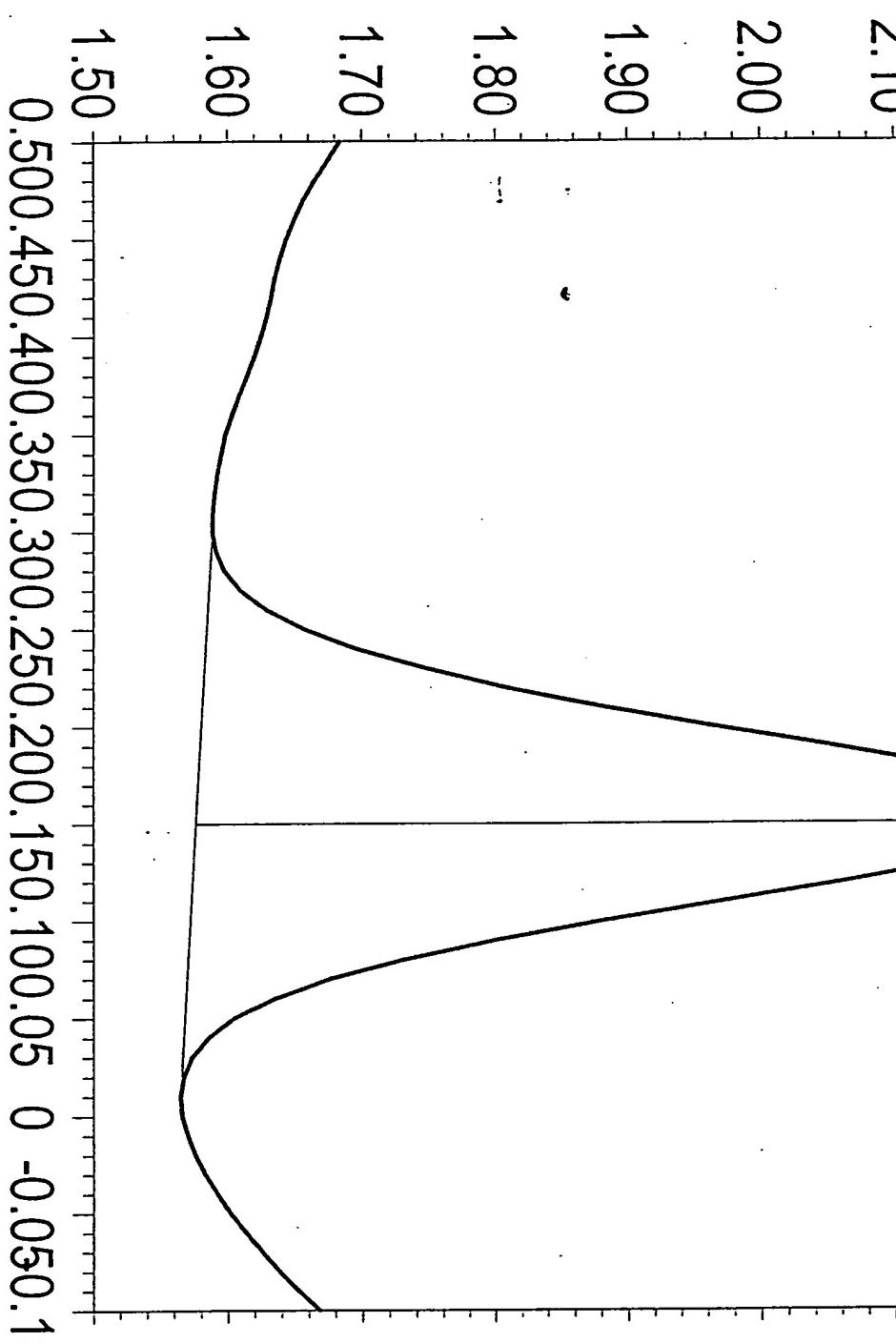
CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

# Electrode #63

Apr. 3, 1998 18:02:37  
Tech: ACV  
File: g200\_033

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

$E_p = 0.150V$   
 $i_p = 6.637e-8A$   
 $A_p = 7.335e-9VA$



CONFIDENTIAL PROPERTY OF  
Clinical Micro Sensors, Inc.

# Electrode #25

May 21, 1998 15:52:41  
Tech: ACV  
File: a367\_007

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7  
 $E_p = 0.140V$   
 $i_p = 2.877e-9A$   
 $A_p = 2.056e-10VA$



Fig.  
20 N

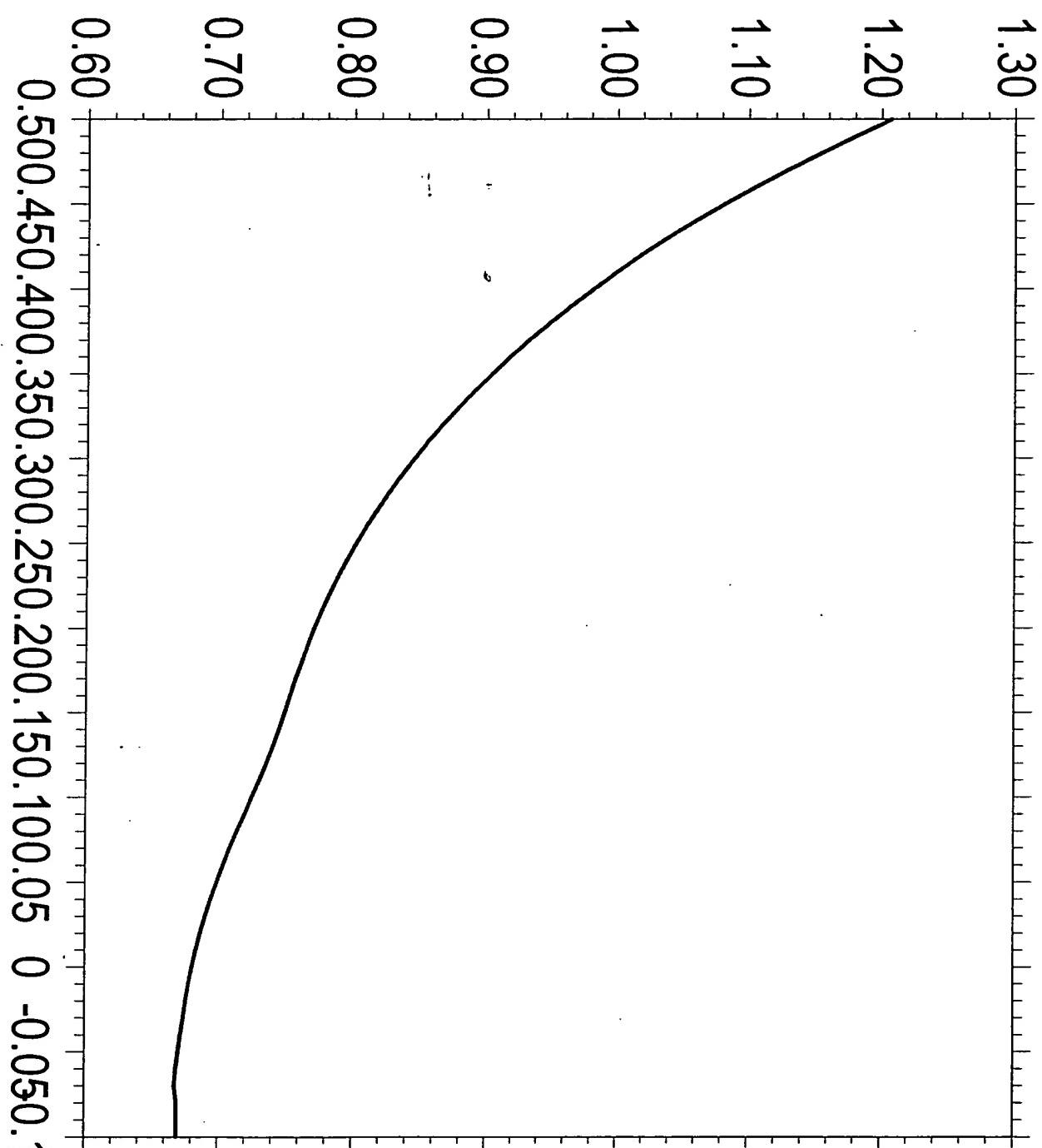
# Electrode #63

May 21, 1998 16:44:35

Tech: ACV  
File: a367\_020

Init E (V) = -0.11  
Final E (V) = 0.5  
Incr E (V) = 0.01  
Amplitude (V) = 0.025  
Frequency (Hz) = 10  
Sample Period (s) = 1  
Quiet Time (s) = 2  
Sensitivity (A/V) = 2e-7

## AC Current / $10^{-7}$ A



Potential / V vs Ag/AgCl

F16-20 C

Sequences for Ligation Experiment

D456

5' - (N6)G(N6) CT(N6) C(N6)G (N6)C(N6) TTC TGC ACC GTA GCC ATG AAA GAT TGT ACT GAG - 3'

D368

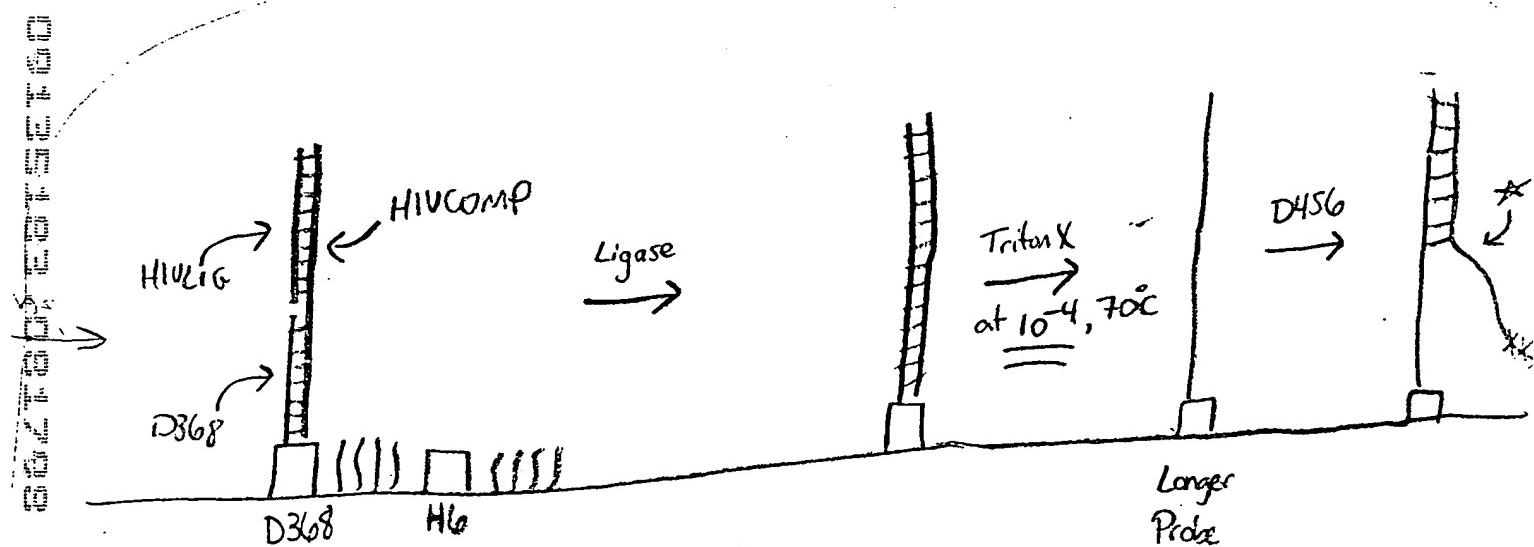
5' - (H2)CC TTC CTT TCC ACA U - 4 UNIT WIRE (C11) - 3'

HIVCOMP

5' - ATG TGG AAA GGA AGG ACA CCA AAT GAA AGA TTG TAC TGA GAG ACA GGC TAA TTT TTT AGG  
GAA GAT CTG G - 3'

HVLIG

5' - CCA GAT CTT CCC TAA AAA ATT AGC CTG TCT CTC AGT ACA ATC TTT CAT TTG GTG T - 3'



$$\text{surface} = \text{D368}/\text{H6}/\text{HIVLIG}$$

\* this detachment point is above the ligation point,  
so that a surface probe that was not ligated would  
not signal.

FIG.  
21

Old 8-Fc, 20-Fc, and 40-Fc signalling molecules

Other ip's

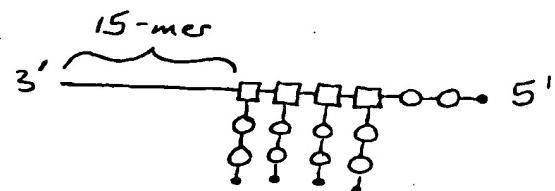
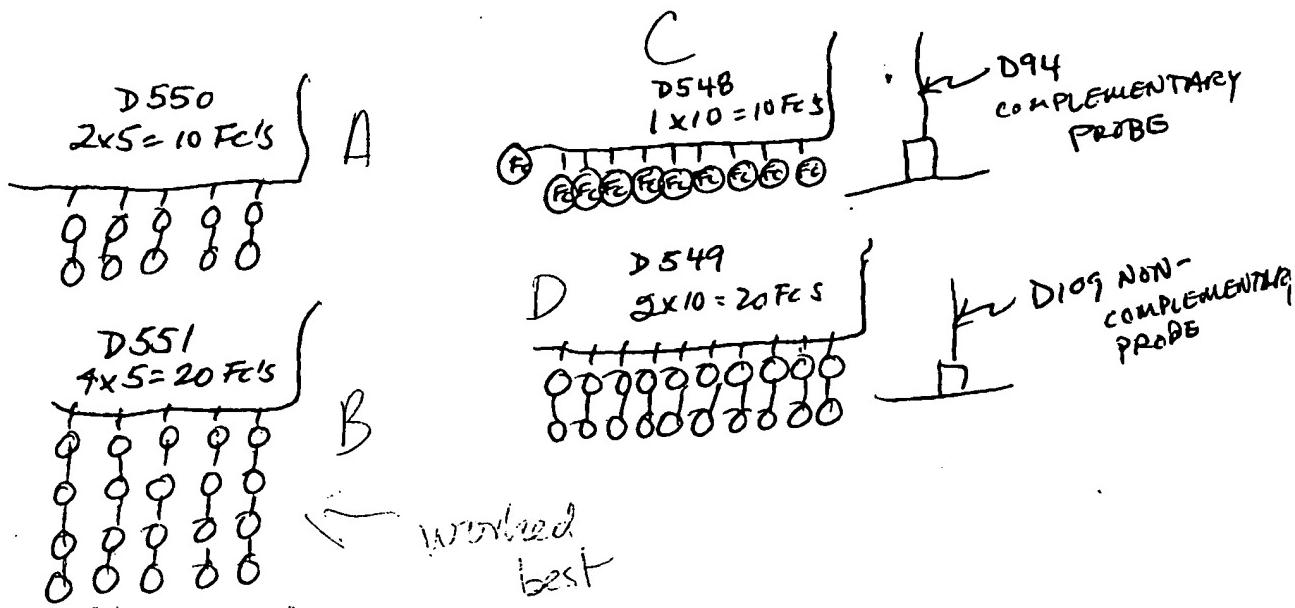
Figs  
23A

Measurer	File	Electrode #	Hybrid code	ip (nA)	Average ip (nA)	STD EV ip (nA)	Potential (mV)	ip (nA)	Potential (mV)
A	1	7	16	0	0	—	—	1.593	70
B	3	8	EU2+reg helpers+reg system	0	0.36	0.71	—		
B	4	6		1.42					
JB	3	5							
A	2	3	1+	0.7449	0.196	0.63	0.29	160	
B	1	4	rRNA EU2+reg helpers+reg system	0.8547	0.63	0.29	160		
JB	1	1							
JB	2	2		0.722					
A	5	13	(2) EU2+EU1,2 reg helpers+reg system	0.3146	0.3441	0.19	0.17	170	0.2506
A	6	15		0					0.8442
JB	4	14							80
JB	6	16		0.11					
A	3	11	(2*) rRNA EU2+EU1,2 reg helpers+reg system	0.586	2.661	1.06	0.51	170	0.05
A	4	12		1				160	
B	2	9		1.6				150	2.4
A	8	22	(3) (2) 20-Fc ETMs+reg system	0.9	3.03	2.99	160		
B	5	23		1.2			160	2.8	120
B	8	24							
JB	7	21		7.376					
A	7	18	3+	1.756					
B	6	19	rRNA+(2) 20-Fc ETMs+reg system	0.77	2.99	2.76	170	0.4778	350
B	7	20		7			120		
JB	5	17		2.448			150		
A	11	29	4	1.426					
B	10	32	(2) 40-Fc ETMs+reg system	3	2.42	1.11	180	0.1	70
B	11	31		3.7			150		
JB	9	30		1.571			170		
A	9	25	4+	12.49					
B	10	26	rRNA+(2) 40-Fc ETMs+reg system	9.278	7.46	4.16	160		
B	9	28		4			130		
JB	8	27		4.088			150		

Peaks w/ $E_o < 160$  mV

Measurer	File	Electrode	Hybrid Code	$2/\pi \cdot i_p$ (nA)			$E_o$ (mV)	$2/\pi \cdot i_p$ (nA)	$E_o$ (mV)
				Raw Data	Average	STDEV			
JZ	2	46	5-	1.041	1.93	1.25	170	4.465	60
A	3	47	2.811				170		
A	1	41	5.7				170		
JZ	1	43	5+	1.862	3.39	2.03	170		
A	2	44	2.613				180		
A	5	53	6-	0.6566			170	2.1	60
JZ	5	55	0.8548				170		
A	6	56	5.167				180	1.64	60
JZ	3	49	5.799				170		
A	4	50	8.468				180		
JZ	4	52	3.187				160		
JZ	7	61	0.1988				160	1.147	60
A	8	62	1.382				170	1.04	50
JZ	8	64	0.6104				160	0.1958	60
JZ	6	58	7+	1.459	1.25	0.29	160	2.38	60
A	7	59	1.042				160		
JZ	10	70	8-	0.3208	0.56	0.34	160	0.504	60
A	11	71	0.7994				190	2.22	60
A	9	65	3.297				170		
JZ	9	67	8+	1.492	2.54	0.94	160		
A	10	68	2.841				170	0.71	60
JZ	12	76	9-	1.215	1.22	#DIV/0!	170	4.414	50
JZ	11	73	9+	3.768	4.68	1.29	170	0.7741	50
A	12	74	5.592				170	0.53	60
JZ	14	78	2.842				170	2.319	50
A	14	80	10-	7.4	5.12	3.22	170		
A	13	77	5.582				170		
JZ	13	79	10+	4.337	4.96	0.88	160	3.173	50

FBS  
23B



□ = N38  
 ○ = C23  
 • = H2

E

FIG 24

Fig 24 cont

F  
Bristle-Attached Fc's

$\text{v} = 10 \text{ Hz}$ ,  $\epsilon = 25 \text{ mV}$

measurer	expt	file	electrode	surface	hybrid	$2/\pi \cdot i_p (\text{nA})$	$E_0 (\text{mV})$	average $2/\pi \cdot i_p (\text{nA})$	STDEV $2/\pi \cdot i_p (\text{nA})$
A	409	1	1	<u>"+"</u> surface 2:2:1 D94/H6/M44*, total thiol = 833 uM	D548 (1x10)**	22.6	150	14.5	5.8
A	409	17	17			9.622	200		
Z	73	8	8			14.51	100		
Z	73	22	24			11.15	110		
A	409	8	7		D549 (2x10)	53.52	200	60.6	12.9
A	409	22	23			71.13	220		
Z	73	1	2			71.66	110		
Z	73	17	18			45.9	120		
A	409	4	3		D550 (2x5)	72.4	190	45.5	18.9
A	409	18	19			30.67	210		
Z	73	7	6			44.49	120		
Z	73	19	22			34.43	120		
A	409	7	5		D551 (4x5)	105.8	210	74.9	23.5
A	409	19	21			48.66	230		
Z	73	4	4			70.42	130		
Z	73	18	20			74.77	130		
A	409	9	9		D548 (1x10)	5.665	200	1.6	2.7
A	409	25	25			0.6443	250		
Z	73	16	16			0.0864	120		
Z	73	30	32			0	-		
A	409	16	15		D549 (2x10)	10.24	230	8.3	5.9
A	409	30	31			14.57	260		
Z	73	9	10			7.881	130		
Z	73	25	26			0.5476	140		
A	409	12	11		D550 (2x5)	4.513	230	3.7	1.6
A	409	26	27			4.264	260		
Z	73	15	14			4.553	150		
Z	73	27	30			1.314	140		
A	409	15	13		D551 (4x5)	10.31	240	9.0	6.9
A	409	27	29			17.46	280		
Z	73	12	12			7.445	160		
Z	73	26	28			0.8812	90		

Note: M44 = M43. \*\* Also note: (n x m) means there are m bristles, each with n Fc's.

ERTY OF  
S. inc

## Base-Replacement Fc's (June 3, 1998); $v = 10$ Hz, $\epsilon = 25$ mV

measurer	expt	file	electrode	surface	hybrid	$2/\pi \cdot i_p$ (nA)	$E_0$ (mV)	averag $2/\pi \cdot i_p$ (nA)	STDEV $2/\pi \cdot i_p$ (nA)
A	52	1	1	"+" surface	10 uM D405	4.81	170		
A	52	4	3	2:2:1 D94/H6/M44*, total thiol = 833 uM	In 6x SSC w/50% FCS	20.63	180		
Z	384	1	2			37.42	170	18.04	14.53
Z	384	4	4			9.31	160		
A	52	7	5	"-" surface	10 uM D405	0.1	160		
A	52	10	7	2:2:1 D109/H6/M44*, total thiol = 833 uM	In 6x SSC w/50% FCS	9.97	160		
Z	384	5	6			0	--	3.12	4.70
Z	384	8	8			2.425	180		

\* Note: M44 = M43.

B

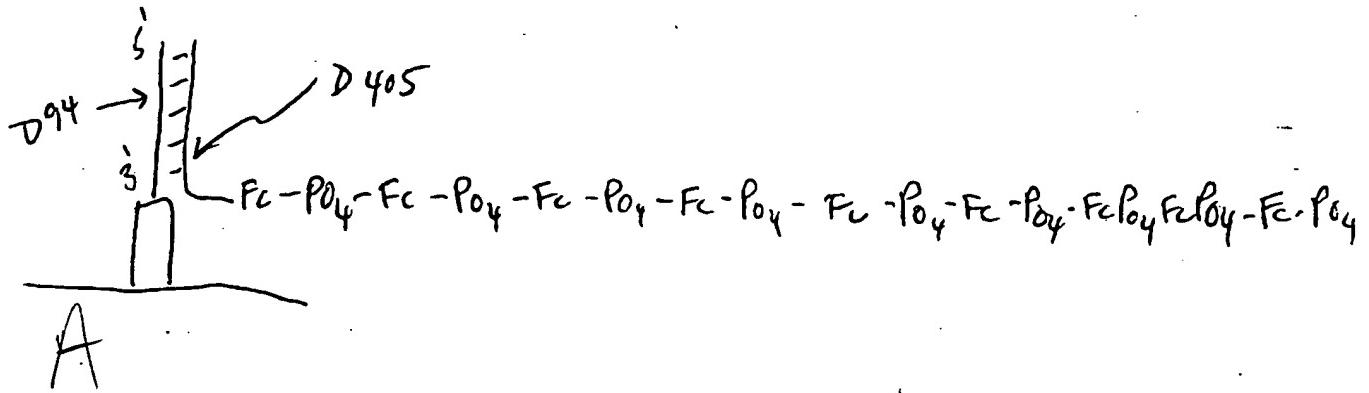
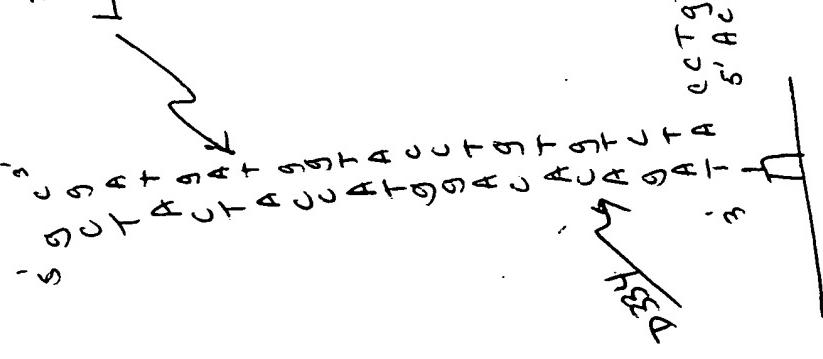


Fig 25

P<sup>4</sup>  
expt

P = positive

LP 280



5' G C T A + G A T C U G A T G A C U T G A T C U T A G T C T G T C T C A T T A G T C T G T C \* G T C \* G T C \* T G G \* T 3'  
→ 335

Fig.  
26A

\* N6 Fc

Fig.  
263

**PCR Amplification Monitored by Electronic Sensing  
for Differing Initial Numbers of Template**

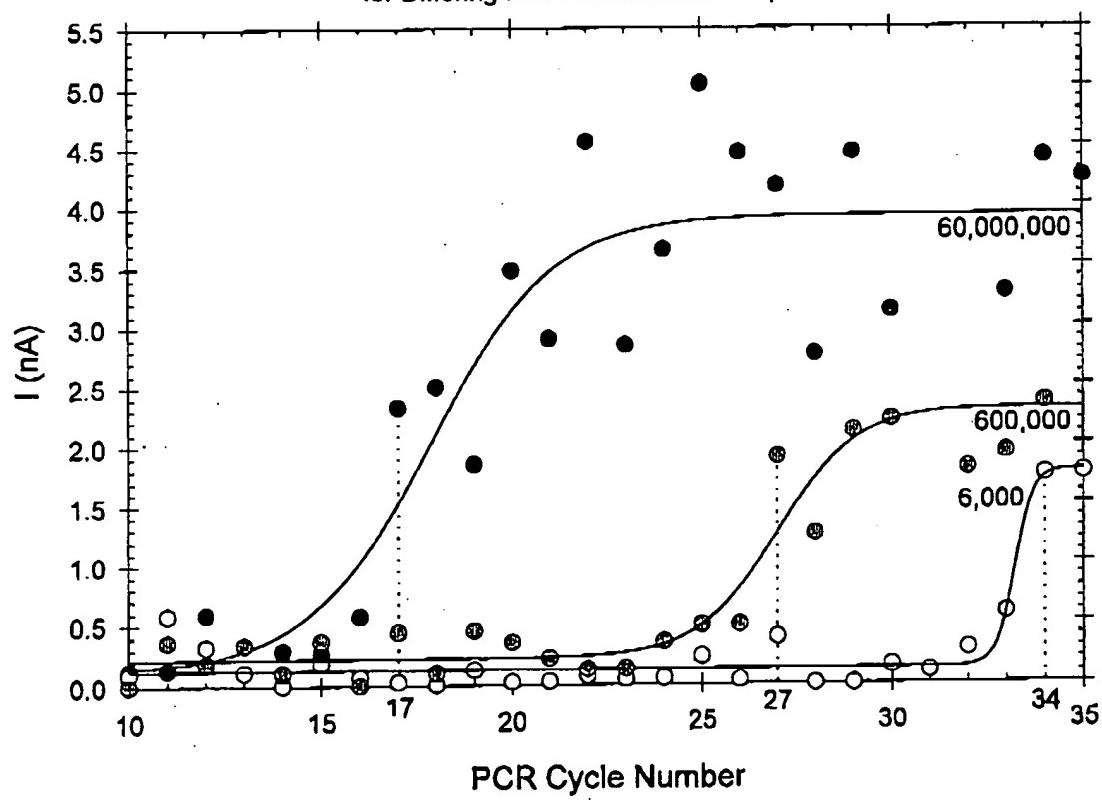


Fig 27